1/2/15/16/10

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NEWS
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NEWS
         MAR 15
                 WPIDS/WPIX enhanced with new FRAGHITSTR display format
NEWS
      3
         MAR 16
                 CASREACT coverage extended
NEWS
         MAR 20
                 MARPAT now updated daily
NEWS
         MAR 22 LWPI reloaded
NEWS · 6
         MAR 30
                 RDISCLOSURE reloaded with enhancements
NEWS
         APR 02
                 JICST-EPLUS removed from database clusters and STN
NEWS
         APR 30
     8
                 GENBANK reloaded and enhanced with Genome Project ID field
NEWS
     9
         APR 30
                 CHEMCATS enhanced with 1.2 million new records
NEWS 10
         APR 30
                 CA/CAplus enhanced with 1870-1889 U.S. patent records
NEWS 11
         APR 30
                 INPADOC replaced by INPADOCDB on STN
NEWS 12
         MAY 01
                 New CAS web site launched
NEWS 13
         MAY 08
                 CA/CAplus Indian patent publication number format defined
NEWS 14
         MAY 14
                 RDISCLOSURE on STN Easy enhanced with new search and display
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NEWS 15
                 BIOSIS reloaded and enhanced with archival data
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NEWS 17
         MAY 21
                 CA/CAplus enhanced with additional kind codes for German
                 patents
NEWS 18
         MAY 22
                 CA/CAplus enhanced with IPC reclassification in Japanese
                 patents
NEWS 19
         JUN 18
                 CA/CAplus to be enhanced with pre-1967 CAS Registry Numbers
              NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT
              MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
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1565,113 6/26/2007 6/26/2007

NEWS HOURS STN Operating Hours Plus Help Desk Availability
NEWS LOGIN Welcome Banner and News Items
NEWS IPC8 For general information regarding STN implementation of IPC 8

AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.

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for Connie

=> file reg
COST IN U.S. DOLLARS

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ENTRY SESSION
0.21 0.21

FULL ESTIMATED COST

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http://www.cas.org/support/stngen/stndoc/properties.html

=> s allylamine

L1 1332 ALLYLAMINE

=> s l1 and poly 1839686 POLY

L2 505 L1 AND POLY

=> s methylcarboxy and 12

19 METHYLCARBOXY

L3 · 0 METHYLCARBOXY AND L2

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 15.30 15.51

FULL ESTIMATED COST '

FILE 'CAPLUS' ENTERED AT 18:45:45 ON 26 JUN 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 26 Jun 2007 VOL 147 ISS 1 FILE LAST UPDATED: 25 Jun 2007 (20070625/ED)
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233684 PROTECTIVE

L4 3094 AMINE AND PROTECTIVE

=> s 14 and allyl 106612 ALLYL

L5 109 L4 AND ALLYL

=> s 15 and poly 705041 POLY

L6 3 L5 AND POLY

=> d all 1-3

```
L6 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN
```

AN 2004:252464 CAPLUS

DN 140:255816.

ED Entered STN: 26 Mar 2004

TI Multistage gas generator with multiple chambers, propellants, and separate initiation

IN Daoud, Sami

PA Textron Systems Corporation, USA

SO PCT Int. Appl., 47 pp. CODEN: PIXXD2

DT Patent

LA English

IC ICM C06D005-06

ICS F42B012-58; F42B010-56; B60R021-26

CC 50-1 (Propellants and Explosives)

FAN. CNT 1

| FAN. | PATENT NO. | | | | | KIND | | DATE | | | APPLICATION NO. | | | | | DATE | | | |
|------|------------|--------------------------|------------|------------|------------|------------|------------|-------------------|------------|------------|-----------------|------------|------------|------------|------------|----------|----------|-----|--|
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| | | W: | CO, | CR, | CU, | CZ, | DE, | AU,
DK,
IL, | DM, | DZ, | EC, | EE, | EG, | ES, | FI, | GB, | GD, | GE, | |
| | | • | OM, | PG, | PH, | PL, | PT, | MA,
RO,
UG, | RU, | SC, | SD, | SE, | SG, | SK, | SL, | | | | |
| | | RW: | GH,
KG, | GM,
KZ, | KE,
MD, | LS,
RU, | MW,
TJ, | MZ,
TM,
IE, | SD,
AT, | SL,
BE, | SZ,
BG, | TZ,
CH, | UG,
CY, | ZM,
CZ, | ZW,
DE, | DK, | EE, | ES, | |
| | US | BF, BJ, CF, 2004050283 | | | CG, | CI, | CM, | GA, | GN, | GQ, | G₩, | ML, | MR, | ŅΕ, | SN, | TD, | TG | | |
| | US | 2004056458 | | | | A1 | | | 0325 | į | US 2002-242494 | | | | | 20020912 | | | |
| | | 6877435
2003270501 | | | | | | | | | AU 2003-270501 | | | | | 20030910 | | | |

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A2
                                           EP 2003-752198
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                                                                    20030910
                                             JP 2004-536442
PRAI US 2002-242494
                          Α
                                20020912
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                                20020912
     WO 2003-US28373
                                20030910
CLASS
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                        C06D005-06
                 ICS
                        F42B012-58; F42B010-56; B60R021-26
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                        C06B0031-00 [ICS,7,C*]; C06D0005-00 [ICS,7]
                        3D054/DD17; 3D054/DD21; 3D054/DD22; 3D054/DD28;
                 FTERM
                        3D054/FF15; 3D054/FF18; 4G068/DA08; 4G068/DA10;
                        4G068/DB10; 4G068/DB30; 4G068/DD15
AΒ
     A multichamber gas generator unit is characterized by having two or more
     compartments, each with a sep. initiator, in which the propellants may
     have different geometries, can be initiated at different times, and can
     have different rates of gas evolution. The compns. can be varied to
     tailor different rates of gas production (e.g., to produce a rapid initial
     inflation, followed by a more gradual inflation rate in subsequent
     stages). Propellant compns. can be comprised of an oxidizer 84-95, a fuel
     3.4-13.4, and a binder 1.5-2.6 weight%, and can include: (1) fuels, such as
     CL-20, RDX, HMX, GAP, NGU, TATB, LLM-105, and EDNA, (2) binders, such as
     polycaprolactone, polyisobutene, glycidyl azide polymer, and poly
     (vinylpyrrolidone), and ammonium nitrate oxidizer. Addnl. components
     include a flash suppressant (e.g., potassium nitrate), stabilizers (Et
     centralite, Acardite, diphenylamine, etc.), combustion inhibitors for
     control of ballistic spikes and an inordinately high initial pressure
     rise, d.-adjusting agents, static dissipators, etc.
ST
     gas generator propellant multiple chamber initiation; nitramine fuel gas
     generator propellant multiple initiation
ΙT
     Polyesters, uses
     RL: NUU (Other use, unclassified); TEM (Technical or engineered material
     use); USES (Uses)
        (combustion inhibitors and binders; multistage gas generator with
        multiple chambers, propellants, and sep. initiation)
ΙT
     Propellants (fuels)
        (composite, gas-generating; multistage gas generator with multiple
        chambers, propellants, and sep. initiation)
ΙT
     Polyoxyalkylenes, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (energetic, binders; multistage gas generator with multiple chambers,
        propellants, and sep. initiation)
ΙT
     Airbags (protective)
        (inflation of; multistage gas generator with multiple chambers,
        propellants, and sep. initiation)
ΙT
     Accelerometers
        (microaccelerometers, gas generators for; multistage gas generator with
        multiple chambers, propellants, and sep. initiation)
IT
     Gas generators
        (multistage gas generator with multiple chambers, propellants, and sep.
        initiation)
ΙT
     Nitramines
     RL: TEM (Technical or engineered material use); USES (Uses)
        (propellants; multistage gas generator with multiple chambers,
        propellants, and sep. initiation)
ΙT
     Ammunition
        (tailored propellants for; multistage gas generator with multiple
        chambers, propellants, and sep. initiation)
```

```
9003-27-4, Polyisobutene 9003-39-8, Polyvinylpyrrolidone
                                                                  24980-41-4.
     Polycaprolactone 25248-42-4, Polycaprolactone
     RL: NUU (Other use, unclassified); TEM (Technical or engineered material
     use); USES (Uses)
        (binder; multistage gas generator with multiple chambers, propellants,
        and sep. initiation)
     96-05-9, Allyl methacrylate 97-90-5, Ethylene glycol
IT
     dimethacrylate
                    2358-84-1, Diethylene glycol dimethacrylate
     RL: NUU (Other use, unclassified); TEM (Technical or engineered material
     use); USES (Uses)
        (combustion inhibitors; multistage gas generator with multiple
        chambers, propellants, and sep. initiation)
     7757-79-1, Potassium nitrate, uses
IΤ
     RL: NUU (Other use, unclassified); TEM (Technical or engineered material
     use); USES (Uses).
        (flash suppressant; multistage gas generator with multiple chambers,
        propellants, and sep. initiation)
IT
     6484-52-2, Ammonium nitrate, uses
     RL: NUU (Other use, unclassified); TEM (Technical or engineered material
     use); USES (Uses)
        (oxidizer; multistage gas generator with multiple chambers,
        propellants, and sep. initiation)
    121-82-4, RDX
                    505-71-5, EDNA
                                      2691-41-0, HMX
                                                       3058-38-6, TATB
                                    132683-64-8 135285-90-4, CL-20
     55510-04-8, Dinitroglycoluril
     143178-24-9, Glycidyl azide polymer
     RL: TEM (Technical or engineered material use); USES (Uses)
        (propellants; multistage gas generator with multiple chambers,
        propellants, and sep. initiation)
     85-98-3, Ethyl centralite 119-75-5, 2-Nitrodiphenyl amine
IT
     122-39-4, Diphenyl amine, uses 603-54-3, Akardite I
     13114-72-2, Akardite II
     RL: NUU (Other use, unclassified); TEM (Technical or engineered material
     use); USES (Uses)
        (stabilizer; multistage gas generator with multiple chambers,
       'propellants, and sep. initiation)
ΙT
     7782-42-5, Graphite, uses
     RL: NUU (Other use, unclassified); TEM (Technical or engineered material
     use); USES (Uses)
        (static dissipators; multistage gas generator with multiple chambers,
        propellants, and sep. initiation)
L6
     ANSWER 2 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN
AN
     1996:761733 CAPLUS
DN
     126:39398
     Entered STN: 01 Jan 1997
ΕD
     Modified poly(maleic anhydride-co-styrene) and related polymers
TΙ
     as a base for color filters
     Pfeiffer, Michael J.
ΙN
     Brewer Science, Inc., USA
PA
SO
     PCT Int. Appl., 62 pp.
     CODEN: PIXXD2
DT
     Patent
     English
I.A
IC
     ICM G03F009-00 .
     ICS C09K019-00; G03C005-00; G03C005-56
     73-11 (Optical, Electron, and Mass Spectroscopy and Other Related
     Properties)
     Section cross-reference(s): 38, 42, 74
FAN.CNT 1
     PATENT NO.
                         KIND
                                DATE
                                          APPLICATION NO.
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A1 19961024 WO 1996-US5192
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         RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML,
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     AU 9656641
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                         C09K019-00; G03C005-00; G03C005-56
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                         [I,C*]; G02B0005-20 [I,A]; G03F0007-00 [I,C*];
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                         G03F0007-00 [I,A]; G03F0007-00 [I,C*]; G03F0007-038
                         [I,A]; G03F0007-038 [I,C*]
     Methods of producing color filter elements by photolithog. patterning
AB
     entail using a nonparticulate filter coating material to form a layer
     which is patterned and cured; the coating material comprises a vehicle and
     a heat stable soluble dye. The vehicle may include a modified poly
     (maleic anhydride-co-styrene) and related polymers. The polymers may be
     used as vehicles in a clear or partially clear form, as protective
     or blocking and anti-leach/stain coatings.
ST
     color filter manuf polymer base; maleic anhydride styrene polymer filter
     base
IT
     Coating materials
     Dyes
     Lithography
     Optical filters
        (modified polymers as a base for color filters)
     1330-38-7, Solvent Blue 38 12226-78-7, Solvent Blue 67 12227-67-7,
IT
     Solvent Yellow 82 12237-27-3, Solvent Red 119 37229-23-5, Solvent Blue
          61969-48-0, Solvent Red 127
     RL: DEV (Device component use); USES (Uses)
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(modified polymers as a base for color filters)
    107-11-9DP, 2-Propen-1-amine, reaction products with maleic anhydride-styrene copolymer 107-18-6DP, Allyl alcohol, reaction products with maleic anhydride-styrene copolymer 9011-13-6DP,
IT
     Maleic anhydride-styrene polymer, reaction products with alcs. and amines
     25265-97-8P 184530-45-8P 184530-46-9P
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process);
     USES (Uses)
        (modified polymers as a base for color filters)
     ANSWER 3 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN 1987:626067 CAPLUS.
L6.
ΑN
DN
     107:226067
     Entered STN: 12 Dec 1987
ĔD
ΤI
     Optical information recording material
ΙN
     Sato, Tsutomu
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     Ricoh Co., Ltd., Japan
SO
     Jpn. Kokai Tokkyo Koho, 5 pp.
     CODEN: JKXXAF
DT
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     ICS G11B007-24
     74-12 (Radiation Chemistry, Photochemistry, and Photographic and Other
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     Reprographic Processes)
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                               19850826
CLASS
                CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
                _____
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                 ICM B41M005-26
 JP 62046680
                 ICS
                        G11B007-24
                 IPCI
                        B41M0005-26 [ICM, 4]; G11B0007-24 [ICS, 4]
                 IPCR
                        G11B0007-24 [I,C*]; G11B0007-244 [I,A]
                 ECLA
                        G11B007/244
GI.
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AB In the title material made of a support bearing thin recording layers containing an organic dye, ≥1 layer selected from a recording layer, an undercoat layer for the former, and a surface protective layer contains a compound of the formula (Z:NZ1NRR1)n+ Xn- [Z = a group forming a (substituted) cyclic ketone which may be condensed with an aromatic group; R, R1 = (substituted) alkyl, aryl; X = acid anion; n = 0, 1, 2; Z1 = I or II(when m = 2) which may be substituted by allyl, alkoxy, halo, The material, for use in semiconductor laser recording, has high preservation stability toward ambient heat and light. Thus, III and a cyanine dye IV were mixed together at a weight ratio of 100:15 and spin-coated on a poly(Me methacrylate) plate to form a recording layer with a thickness of ca. 500 Å. When the material was used for recording and reading with a 790-nm semiconductor laser beam at a recording frequency of 0.7 MHz and a scanning rate of 1.2 m/s the carrier-to-noise ratio was 55 dB, which did not lower much after exposure to light.

ST optical recording material lightfastness improvement; phenylenediamine deriv optical recording; amine phenylenedi deriv optical recording

IT Recording materials

(optical, containing phenylenediamine derivative, for improved lightfastness)

IT 23178-67-8 26752-32-9 111341-21-0 111341-25-4 111341-26-5

111341-27-6 111341-29-8 111408-84-5

RL: TEM (Technical or engineered material use); USES (Uses) (optical recording material containing, with improved lightfastness)

=> s polyallylamine L7 1966 POLYALLYLAMINE

=> s 17 and protected 114788 PROTECTED L8 19 L7 AND PROTECTED

for Connie

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=> d all 1-19
    ANSWER 1 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
L8
    2007:259942 CAPLUS
ΑN
    146:296424
DN
    Entered STN: 09 Mar 2007
ED
TT
     Polymeric molecular receptors as phosphate sequestrants
ΙN
    Huval, Chad C.; Holmes-Farley, Stephen Randall; Dhal, Pradeep K.
PA
    Genzyme Corporation, USA
SO
     PCT Int. Appl., 136pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
IC
     ICM A61K
     35-8 (Chemistry of Synthetic High Polymers)
CC
     Section cross-reference(s): 63
FAN.CNT 1
                                            APPLICATION NO.
                                                                   DATE
     PATENT NO.
                                DATE
                         KIND
                                           _____
     ______
                        ____
                               _______
    WO 2007027566
                                20070308
                                           WO 2006-US33437
                         A2
                                                                  20060825
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
             GE, GH, GM, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP,
             KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN,
             MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS,
             RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
         RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
             IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,
             CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
             GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM
PRAI US 2005-713991P
                         Р
                                20050902
     US 2005-734462P
                                20051108
                          Ρ.
CLASS
                 CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
WO 2007027566
                 ICM
                        A61K
                 IPCI
                        A61K [ICM, 7:]
     Polymers and compns. utilizing such polymers are disclosed for treating
AB
     hyperphosphatemia and other illnesses associated with elevated serum
     phosphate levels. Phosphate binding polymers, or a pharmaceutically
     acceptable salt of the polymers, comprise pendent groups extending from a
     backbone of the polymer. Each pendent group comprises at least two
     nitrogen-bearing functional groups which bind phosphate. A typical
     polymer was manufactured by reaction of 10.2 g polyepichlorohydrin 48 h at
     180° with 200 mL tris(2-aminoethyl)amine in 250 mL NMP.
     phosphate sequestrant amine side group polymer; polyepichlorohydrin
ST
     trisaminoethylamine adduct manuf
IT
     Polyethers, preparation
     Polyolefins
     Polyoxyalkylenes, preparation
     RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological
     study); PREP (Preparation); USES (Uses)
        (amine derivs.; polymeric mol. receptors having amine side chains as
        phosphate sequestrants)
ΙT
     Kidney, disease
        (hyperphosphatemia; polymeric mol. receptors having amine side chains
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as phosphate sequestrants)

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IT
     Sequestering agents
        (polymeric mol. receptors having amine side chains as phosphate
        sequestrants)
ΙT
     Phosphates, processes
     RL: REM (Removal or disposal); PROC (Process)
        (polymeric mol. receptors having amine side chains as phosphate
        sequestrants)
ΙT
     Quaternary ammonium compounds, preparation
     RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological
     study); PREP (Preparation); USES (Uses)
        (polymers; polymeric mol. receptors having amine side chains as
        phosphate sequestrants)
IT
     56-18-8DP, N-(3-Aminopropyl)-1,3-propanediamine, reaction products with
     polyepichlorohydrin
                           107-15-3DP, Ethylenediamine, reaction products with
     polyepichlorohydrin
                           111-40-0DP, Diethylenetriamine, reaction products
     with polyepichlorohydrin
                                4097-89-6DP, Tris(2-aminoethyl)amine, reaction
     products with polyepichlorohydrin
                                         24969-06-ODP, Polyepichlorohydrin,
     reaction products with polyamines
                                         29160-08-5DP, Poly[(2-chloroethyl)
     vinyl ether], reaction products with tris(aminoethyl)amine
                                                                  32610-74-5P,
     4-[[Bis(2-aminoethyl)amino]methyl]styrene homopolymer
                                                             61710-61-0DP,
     Polyepichlorohydrin, sru, reaction products with polyamines
     928164-64-1P, N-[2-[Bis(2-aminoethyl)amino]ethyl]methacrylamide
                   928164-68-5P, 4-[[Bis(3-aminopropyl)amino]methyl]styrene
     homopolymer
                   928164-72-1P, N-[3-(4-Vinylbenzylamino)propyl]-1,3-
     homopolymer
     propanediamine homopolymer
     RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical
     process); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); PROC (Process); USES (Uses)
        (polymeric mol. receptors having amine side chains as phosphate
        sequestrants)
                                                  25014-12-4DP,
     9003-05-8DP, Polyacrylamide, amine derivs.
ΙT
     Polymethacrylamide, amine derivs. 25014-13-5P, Epichlorohydrin-
     ethylenediamine copolymer
                                 25085-17-0P, Diethylenetriamine-
                                 26336-38-9P, Polyvinylamine
                                                               30551-89-4P,
     epichlorohydrin copolymer
     Polyallylamine
                      37017-08-6DP, N-Acryloyloxysuccinimide
     homopolymer, reaction products with BOC-partially protected
                             41077-50-3DP, 3,7,9-Triazabicyclo[3.3.1]nonane,
     polyamine, deprotected
                       51961-45-6P, N-(3-Aminopropyl)-1,3-propanediamine-
     polymer derivs.
     epichlorohydrin copolymer
                                 113506-17-5P, Epichlorohydrin-tris(2-
     aminoethyl) amine copolymer
                                  928164-61-8P, (2-Chloroethyl) vinyl
     ether-epichlorohydrin-tris(2-aminoethyl)amine copolymer
                                                               928164-65-2P,
     N-[2-[Bis(2-aminoethyl)amino]ethyl]methacrylamide-epichlorohydrin
                 928164-66-3P, 4-[[Bis(2-aminoethyl)amino]methyl]styrene-
     copolymer '
                                 928164-69-6P, 4-[[Bis(3-
     epichlorohydrin copolymer
     aminopropyl)amino]methyl]styrene-epichlorohydrin copolymer
     Epichlorohydrin-N-[3-(4-vinylbenzylamino)propyl]-1,3-propanediamine
                 928164-76-5P, 4-[[Bis(2-aminoethyl)amino]methyl]styrene-N, N'-
     copolymer
     ethylenebisacrylamide copolymer 928164-78-7P, 4-[[Bis(2-
     aminoethyl)amino]methyl]styrene-N, N'-bis(4-vinylbenzyl)ethylenediamine
                928164-85-6DP, reaction products with N-Acryloyloxysuccinimide
     copolymer
     homopolymer, deprotected 928164-88-9P, N, N-Bis(2-aminoethyl)-N', N'-
                                          928164-89-0DP, 2,4-
     diallylethylenediamine homopolymer
     Bis(aminomethyl)piperidine, polymer derivs.
                                                  928164-90-3DP,
     2,5-Bis(aminomethyl)tetrahydropyrrole, polymer derivs.
     RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological
     study); PREP (Preparation); USES (Uses)
        (polymeric mol. receptors having amine side chains as phosphate
        sequestrants)
     107-15-3, 1,2-Ethylenediamine, reactions
ΙT
     RL: RCT (Reactant); RACT (Reactant or reagent)
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(polymeric mol. receptors having amine side chains as phosphate
        sequestrants)
ΙT
     7778-77-0, Potassium dihydrogen phosphate
     RL: REM (Removal or disposal); PROC (Process)
        (polymeric mol. receptors having amine side chains as phosphate
        sequestrants)
     29160-08-5P, Poly[(2-chloroethyl) vinyl ether] 46734-05-8P,
     4-[[Bis(2-aminoethyl)amino]methyl]styrene
                                                161038-11-5P,
     4-(2-Aminoethyl)-1,7-bis(tert-butoxycarbonyl)-1,4,7-triazaheptane
     884862-51-5P, N,N'-Bis(4-vinylbenzyl)ethylenediamine
                                                            928164-62-9P
     928164-63-0P, N-[2-[Bis(2-aminoethyl)amino]ethyl]methacrylamide
                                                                928164-70-9P.
     928164-67-4P, 4-[[Bis(3-aminopropyl)amino]methyl]styrene
     N-[3-(4-Vinylbenzylamino)propyl]-1,3-propanediamine
                                                           928164-81-2P
     928164-83-4P 928164-85-6P
                                   928164-87-8P, N, N-Bis(2-aminoethyl)-N', N'-
     diallylethylenediamine
     RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT
     (Reactant or reagent)
        (precursor; polymeric mol. receptors having amine side chains as
        phosphate sequestrants)
     56-18-8, Bis(3-aminopropyl)amine
                                      107-13-1, Acrylonitrile, reactions
TΤ
     920-46-7, Methacryloyl chloride 1592-20-7, 4-Vinylbenzyl chloride
     4097-89-6, Tris(2-aminoethyl)amine
                                          4963-47-7, Tris(3-aminopropyl)amine
     24424-99-5, Di-tert-butyl dicarbonate
                                             58632-95-4
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (precursor; polymeric mol. receptors having amine side chains as
        phosphate sequestrants)
     ANSWER 2 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
L8
ΑN
     2007:15155 CAPLUS
DN
     146:322995
     Entered STN: 05 Jan 2007
ED
     DNA-polycation complexes Effect of polycation structure on
ΤI
     physico-chemical and biological properties
UΑ
     Slita, A. V.; Kasyanenko, N. A.; Nazarova, O. V.; Gavrilova, I. I.;
     Eropkina, E. M.; Sirotkin, A. K.; Smirnova, T. D.; Kiselev, O. I.;
     Panarin, E. F.
     Research Institute of Influenza, RAMS, St. Petersburg, 197376, Russia
CS
     Journal of Biotechnology (2007), 127(4), 679-693
SO
     CODEN: JBITD4; ISSN: 0168-1656
PB
     Elsevier B.V.
DT
     Journal
LA
     English
CC
     63-5 (Pharmaceuticals)
     Section cross-reference(s): 35
     The purpose of the study was to investigate the influence of cationic
AB
     polymer structure on the formation of DNA-polycation complexes and their
     transfection activity. Primary, tertiary, and quaternary polyamines with
     mol. masses ranging from 8000 to 200,000 were investigated. DNA-cationic
     polymer interaction was characterized by low gradient viscometry, dynamic
     light scattering, CD, UV spectrometry, flow birefringence, DNA
     electrophoresis, and electron microscopy. Transfection activity of the
     complexes was evaluated by the expression of reporter gene
     (\beta-galactosidase) and using synthetic FITC-labeled oligonucleotides.
     Complex formation was found to be dependent on the structure and mol. weight
     of the polymer and the ionic strength of the solution Secondary DNA
     structure in complexes was not disrupted, and DNA was protected
     from protonation. Cell lines of different origin were used for testing of
     transfection activity of the complexes. The sensitivity of the cells to
     transfection was established to be highly dependent on the cell line.
     DNA-polycation complexes are non-toxic according to MTT.
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ST

ΙT

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RE

Polyallylamine, and polydimethylaminoethylmethacrylate were found to be the most promising polycations for gene delivery. Transfection efficacy of their complexes with DNA to T-98G cells reaches up to 90-100%. It was found that optimal mol. mass of polydimethylaminoethylmethacrylate is in the range of 8000-50,000 Da. polycation DNA polyallylamine polydimethylaminoethylmethacrylate transfection gene vector Genetic vectors Molecular association Particle size Viscosity (DNA-polycation complexes Effect of polycation structure on physico-chemical and biol. properties) DNA RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (DNA-polycation complexes Effect of polycation structure on physico-chemical and biol. properties) 25104-18-1DP, Poly(L-lysine), complex with DNA 25154-86-3DP, Polydimethylaminoethylmethacrylate, complex with DNA 26161-33-1DP, complex with DNA 29792-49-2DP, Polyvinylamine hydrochloride, complex 38000-06-5DP, Poly(L-lysine), complex with DNA with DNA 71550-12-4DP, Polyallylamine hydrochloride, complex with DNA 124221-13-2DP, complex with DNA 124221-14-3P 124221-17-6P RL: ADV (Adverse effect, including toxicity); BSU (Biological study, unclassified); PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (DNA-polycation complexes Effect of polycation structure on physico-chemical and biol. properties) RE.CNT THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS RECORD (1) Akinc, A; J Gene Med 2005, V7, P657 CAPLUS (2) Bos, G; Vaccine 2004, V23, P460 CAPLUS (3) Dash, P; J Control Rel 1997, V48, P269 CAPLUS (4) De Smedt, S; Pharm Res 2000, V17, P113 CAPLUS (5) Dubruel, P; Biomacromolecules 2004, V5, P379 CAPLUS (6) Dubruel, P; Eur J Pharm Sci 2003, V18, P211 CAPLUS (7) Eldred, S; Bioconjug Chem 2005, V16; P694 CAPLUS (8) Frisman, E; Kolloidn Zh 1965, V27, P130 CAPLUS (9) Frisman, E; Mol Biol 1990, V24, P318 CAPLUS (10) Harada, S; Makromol Chem Rapid Commun 1984, V5, P27 CAPLUS (11) Howard, K; Biochim Biophys Acta 2004, V1674, P149 CAPLUS (12) Huey, R; Biopolymers 1981, V20, P2533 CAPLUS (13) Itaka, K; Biomaterials 2003, V24, P4495 CAPLUS (14) Kabanov, A; Bioconjug Chem 1995, V6, P7 CAPLUS (15) Kasyanenko, N; Biophys Chem 1998, V70, P93 CAPLUS (16) Kasyanenko, N; Colloids Surf A: Physicochem Eng Aspects 1999, V148, P121 **CAPLUS** (17) Kasyanenko, N; Langmuir 1999, V15, P7912 CAPLUS (18) Mosmann, T; J Immunol Methods 1983, V65, P55 MEDLINE (19) Panarin, E; Makromol Chem 1985, Suppl 9, P25 (20) Panarin, E; Vysokomolek Soed 1977, V19B, P251 (21) Pavlov, G; Prog Colloid Polym Sci 2006, V131, P134 CAPLUS (22) Pavlov, G; Vysokomolek Soed 2004, V46A, P1732 (23) Reschel, T; J Control Rel 2002, V81, P201 CAPLUS

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(26) Suyber, S; Anal Biochem 1975, V64, P284

(25) Stickler, M; Angew Makromol Chem 1984, V123-124, P85

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11/245136
(27) Thomas, M; Appl Microbiol Biotechnol 2003, V62, P27 CAPLUS
(28) van de Wetering, P; J Control Rel 1997, V49, P59 CAPLUS (29) van de Wetering, P; J Control Rel 1998, V53, P145 CAPLUS
(30) Wolfert, M; Bioconjug Chem 1999, V10, P993 CAPLUS
(31) Zhou, J; Biophys Chem 2004, V107, P273
L8
     ANSWER 3 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
ΑN
     2006:187197 CAPLUS
DN
     144:260880
ED
     Entered STN: 02 Mar 2006
     Orthopedic flexible body part fixing materials comprising support and
     curable resins with their surface partially cured
İΝ
     Matsumoto, Yoshikazu; Kato, Yukihiro
     Alcare Co., Ltd., Japan
Jpn. Kokai Tokkyo Koho, 18 pp.
PΑ
     CODEN: JKXXAF
DT
     Patent
LA
     Japanese
CC
     63-7 (Pharmaceuticals)
FAN.CNT 1
     PATENT NO.
                         KIND
                                DATE
                                            APPLICATION NO.
                                                                    DATE
                     . A
     JP 2006055281
                                 20060302
                                             JP 2004-238593
                                                                    20040818
PT
PRAI JP 2004-238593
                                 20040818
CLASS
 PATENT NO.
                 CLASS PATENT FAMILY CLASSIFICATION CODES
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 _____
 JP 2006055281
                 IPCI A61F0013-04 [I,A]; A61F0005-01 [N,A]; A61F0005-02 [N,A]
                 FTERM 4C098/AA01; 4C098/AA02; 4C098/BB05; 4C098/BB09;
                         4C098/BB11; 4C098/BC02; 4C098/BC46; 4C098/DD20;
                        4C098/DD30
     The materials, e.g. splints, casts, etc., comprise a support such as glass
AΒ
     fiber fabric and a curable resin, wherein the most part of the resin
     remains uncured and the resin at the surface is partially cured, e.g. upon
     sheet when the resin is hydraulic or with a light-transmitting protective
     the material is protected because the resin is partially cured,
     thus preventing transfer of the resin during storage and being handled by
     a bare hand. Thus, a 7-ply laminate of a glass fiber fabric coated with a
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- covering the surface with a curing agent-containing water-permeable protective sheet containing curing agents capable of reacting with the resin. Surface of curable composition containing PEGs, bisphenol-type diols, a polyurethane prepolymer (MDI 47.5, modified MDI 11.5%), a catalyst, an antifoamer, etc., was covered with a PET fabric (porosity 60%) containing PVA and a polypropylene elastomer net. The lowermost glass fiber fabric layer was further laminated with a soft cellular polyurethane foam cushion layer, a water-resistant polypropylene melt-blown nonwoven fabric, and a skin-contact PET nonwoven fabric to give a splint material. The material was stored at 50° for 7 days under loading at 19.6~N/100~cm2 to show no resin transfer to the surface net. The material was sprayed with H2O to finally show 3-point bending strength 685 N.
- ST orthopedic flexible fixing material surface partially cured resin; splint material water curable polyurethane glass fiber fabric laminate
- ΙT Polyurethanes, biological studies
 - RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (acrylic; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)
- IT Carbohydrates, reactions Polyamines

IT

IT

IΤ

IΤ

IT

IT

ΙT

ΙT

ΙT

Polysaccharides, reactions RL: RCT (Reactant); RACT (Reactant or reagent) (crosslinking agents; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet) Polyolefin rubber RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (ethylene, net; protective sheet; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet) Polyester fibers, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (fabrics, nonwoven, covering material; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet) Polyester fibers, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (fabrics, protective sheet; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet) Polyurethanes, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (foams, cushion layer; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet) Immobilization, animal Medical goods (orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet) Polyoxyalkylenes, biological studies RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet) Polyamines RL: RCT (Reactant); RACT (Reactant or reagent) (polyalkylene-, crosslinking agents; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet) Polyurethanes, biological studies RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (polyamine-polyoxyalkylene-; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological

(polyamine-polyurethane-; orthopedic flexible fixing materials such as

Polyoxyalkylenes, biological studies

study); PREP (Preparation); USES (Uses)

splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

IT Polyurethanes, biological studies

RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(polyoxyalkylene-; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

IT Polyamines

RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(polyoxyalkylene-polyurethane-; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

IT Plastic foams

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (polyurethane, cushion layer; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

IT Acrylic polymers, biological studies

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (polyurethane-; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

IT Polyolefin rubber

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(propene, net, uppermost covering material; orthopedic flexible fixing
materials such as splint materials comprising support and water- or
light-curable resins with their surface partially cured upon covering
with curing agent-containing protective sheet)

IT Glass fiber fabrics

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (support; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

IT 629-11-8DP, 1,6-Hexanediol, reaction products with polyethylene glycol, bisphenol-type diol, and MDI-based urethane prepolymer RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(crosslinking agent; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

IT 107-15-3, Ethylenediamine, reactions 9002-89-5, Poly(vinyl alcohol) 9002-98-6 25104-18-1, Polylysine 26336-38-9, Poly(vinylamine) 30551-89-4, Polyallylamine 38000-06-5, Polylysine

RL: RCT (Reactant); RACT (Reactant or reagent)
(crosslinking agent; orthopedic flexible fixing materials such as
splint materials comprising support and water- or light-curable resins
with their surface partially cured upon covering with curing
agent-containing protective sheet)

 splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

ΙT 101-68-8DP, MDI, adducts with bisphenol-type diol and modified MDI, reaction products with polyethylene glycol and crosslinking agents 111-40-0DP, Diethylenetriamine, polymers with acrylic urethane oligomers 111-40-0DP, Diethylenetriamine, reaction and hexanediol diacrylate products with polyethylene glycol, bisphenol-type diol, and MDI-based 9002-89-5DP, Poly(vinyl alcohol), reaction products urethane prepolymer with polyethylene glycol, bisphenol-type diol, and MDI-based urethane prepolymer 9002-98-6DP, reaction products with polyethylene glycol, bisphenol-type diol, and MDI-based urethane prepolymer 13048-33-4DP, 1,6-Hexanediol diacrylate, polymers with acrylic urethane oligomers and diethylenetriamine 25322-68-3DP, reaction products with bisphenol-type diol, MDI-based urethane prepolymer, and crosslinking agents 30551-89-4DP, Polyallylamine, polymers with acrylic urethane oligomers and hexanediol diacrylate RL: IMF (Industrial manufacture); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their

IT 9002-88-4, Polyethylene

protective sheet)

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (rubber, net, protective sheet; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

surface partially cured upon covering with curing agent-containing

IT 9003-07-0, Polypropylene

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (rubber, net, uppermost cover; orthopedic flexible fixing materials such as splint materials comprising support and water- or light-curable resins with their surface partially cured upon covering with curing agent-containing protective sheet)

- L8 ANSWER 4 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
- AN 2005:1260368 CAPLUS
- DN 144:171801
- ED Entered STN: 02 Dec 2005
- TI Chemical, thermal, and ultrasonic stability of hybrid nanoparticles and nanoparticle multilayer films
- AU Isaacs, Steven R.; Choo, Hosun; Ko, Weon-Bae; Shon, Young-Seok
- CS Department of Chemistry, Western Kentucky University, Bowling Green, KY, 42101, USA
- SO Chemistry of Materials (2006), 18(1), 107-114 CODEN: CMATEX; ISSN: 0897-4756
- PB American Chemical Society
- DT Journal
- LA English
- CC 37-6 (Plastics Manufacture and Processing)
 Section cross-reference(s): 38
- AB Gold nanoparticles stabilized by thiolate derived from normal and ω -functionalized alkane- and arene-thiols (HS(CH2)nCH3, HS(CH2)10COOH, HS(CH2)11OH, HSC6H4NH2, HSCH2CH2C6H5, and HSCH2C6H4C(CH3)3) were synthesized. Differences in their chemical and ultrasonic stabilities were studied using UV-vis spectroscopy. The results showed that these stabilities of monolayer-protected nanoparticles were greatly influenced by the structure and functionality of the monolayer surrounding a nanoparticle core. The selective functionalization of the nanoparticle

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allowed us to prepare hybrid nanostructure (nanoparticle multilayer assemblies) with different compns. and functionalities. This paper also presents an investigation of the chemical, thermal, and ultrasonic treatments of these nanoparticle multilayer assemblies. The results suggest that the linkers used to build nanoparticle multilayer films govern the overall stability of hybrid nanostructure. gold nanoparticle thiolate multilayer film thermal chem ultrasonic stability Polymer degradation (acoustic; ultrasonic stability of hybrid nanoparticles and multilayer Chemically resistant materials (chemical stability of hybrid nanoparticles and multilayer films) Glass substrates Hybrid organic-inorganic materials Laminated plastic films Monolayers Nanoparticles Self-assembled monolayers (chemical, thermal, and ultrasonic stability of hybrid nanoparticles and multilayer films) Cluster compounds RL: PRP (Properties) (chemical, thermal, and ultrasonic stability of hybrid nanoparticles and multilayer films) Decomposition kinetics (constant; chemical, thermal, and ultrasonic stability of hybrid nanoparticles and multilayer films) (multilayer; chemical, thermal, and ultrasonic stability of hybrid nanoparticles and multilayer films) Polyamines RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (polyamide-, dendrimers; chemical, thermal, and ultrasonic stability of hybrid nanoparticle's and multilayer films) Dendritic polymers RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (polyamide-polyamines; chemical, thermal, and ultrasonic stability of hybrid nanoparticles and multilayer films) Polyamides, properties RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (polyamine-, dendrimers; chemical, thermal, and ultrasonic stability of hybrid nanoparticles and multilayer films) Polymer degradation (thermal; chemical, thermal, and ultrasonic stability of hybrid nanoparticles and multilayer films) 25704-18-1, Polysodium 4-styrenesulfonate 71550-12-4, Polyallylamine hydrochloride RL: POF (Polymer in formulation); PRP (Properties); USES (Uses) (chemical stability of hybrid nanoparticles and multilayer films) 1310-58-3, Potassium hydroxide, reactions 7647-01-0, Hydrochloric acid, reactions RL: RGT (Reagent); RACT (Reactant or reagent) (chemical stability of hybrid nanoparticles and multilayer films) 13770-18-8, Copper perchlorate RL: CAT (Catalyst use); USES (Uses) (chemical, thermal, and ultrasonic stability of hybrid nanoparticles and multilayer films) 1193-02-8D, reaction products with 1-hexane-thiol coated gold clusters

71310-21-9D, reaction products with 1-hexane-thiol coated gold clusters

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73768-94-2D, 11-Mercaptoundecanol, reaction products with 1-hexane-thiol
     coated gold clusters
     RL: PRP (Properties)
        (chemical, thermal, and ultrasonic stability of hybrid nanoparticles and
        multilayer films)
IT
     16903-35-8, Hydrogen tetrachloroaurate
                                               16940-66-2, Sodium
     tetrahydroborate
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (chemical, thermal, and ultrasonic stability of hybrid nanoparticles and
        multilayer films)
     93376-66-0, PAMAM G-2
IT
     RL: POF (Polymer in formulation); PRP (Properties); USES (Uses)
        (dendritic; chemical stability of hybrid nanoparticles and multilayer
        films)
     110-66-7, 1-Pentanethiol 111-31-9, 1-Hexanethiol 111-88-6, 1-Octanethiol 112-55-0, 1-Dodecanethiol 1455-21-6, 1-Nonanethiol
ΙT
                                     25276-70-4, 1-Pentadecanethiol
     4410-99-5, Benzeneethanethiol
     49543-63-7, 4-tert.-Butyl benzyl-mercaptan
     RL: PRP (Properties)
        (gold cluster compound; chemical, thermal, and ultrasonic stability of
        hybrid nanoparticles and multilayer films)
IT
     7440-57-5, Gold, properties
     RL: PRP (Properties)
        (nanoparticle, clusters; chemical, thermal, and ultrasonic stability of
        hybrid nanoparticles and multilayer films)
RE.CNT 36
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- L8 ANSWER 5 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
- AN 2005:1090325 CAPLUS
- DN 144:23396
- ED Entered STN: 12 Oct 2005
- TI pH-Induced Reversible Conformational and Morphological Regulation of Polyleucine Grafted Polyallylamine Assembly in Solution
- AU Higuchi, Masahiro; Inoue, Takateru; Miyoshi, Hidenori; Kawaguchi, Masami
- CS Department of Chemistry for Materials, Faculty of Engineering, Mie University, Tsu, Mie, 514-8507, Japan
- University, Tsu, Mie, 514-8507, Japan SO Langmuir (2005), 21(24), 11462-11467 CODEN: LANGD5; ISSN: 0743-7463
- PB American Chemical Society
- DT Journal
- LA English
- CC 36-7 (Physical Properties of Synthetic High Polymers)
- AB One of the essential parts in the mol. mechanism of biol. properties is the structural changes of proteins induced by stimuli. An amphiphilic copolymer, poly(L-leucine) grafted polyallylamine as a simple model of proteins, has been prepared by NCA (N-carboxyanhydride) polymerization with free amino groups of polyallylamine as an initiator. Here, we report the pH-induced reversible conformational and morphol. regulation of the amphiphilic copolymer, whose hydrophobic peptide graft chains have no pH-sensitive groups, in an aqueous solution containing 50 vol% trifluoroethanol.

The conformation of the poly(L-leucine) graft chain was found to be strongly pH dependent. Under acidic conditions, where electrostatic repulsion existed between the neighboring protonated amine moieties of the polyallylamine main chain, the rapid aggregation of the poly(L-leucine) graft chains was disturbed, and the peptide graft chains formed a β -sheet structure owing to the intramol. hydrogen bonding among the graft chains. Under this condition, the amphiphilic polymer formed amyloid-like fibrils, and then the fibrils grew into a planer plate composed of staked β -sheets. On the other hand, under basic conditions, the poly(L-leucine) graft chains showed conformational transitions from a β -sheet structure to an α -helical conformation owing to a distortion of the regular arrangement of the peptide graft chains by the conformational change of the polyallylamine main chain, whose amino groups were deprotonated. The conformational transition resulted in a disturbance of the regular sheet assembly of the amphiphilic copolymer and induced morphol. changes to the amorphous globular aggregates. The pH-induced conformational and morphol. changes of the poly(L-leucine) graft polyallylamine were reversible and synchronized with the protonation of the polyallylamine main chain.

- ST polyleucine polyallylamine graft copolymer morphol chain conformation hydrogen bonding
- IT Polymer chains
 - (conformation; preparation and pH-induced reversible conformational and morphol. regulation of polyleucine grafted polyallylamine assembly in solution)
- IT Polyamides, properties
 - RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP (Physical process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)
 - (graft polymers; preparation and pH-induced reversible conformational and

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morphol. regulation of polyleucine grafted polyallylamine
        assembly in solution)
ΙT
     Hydrogen bond
        (intramol.; preparation and pH-induced reversible conformational and
        morphol. regulation of polyleucine grafted polyallylamine
        assembly in solution)
ΙT
     Aggregates
     Circular dichroism
     Conformational transition
     Polymer morphology
        (preparation and pH-induced reversible conformational and morphol.
        regulation of polyleucine grafted polyallylamine assembly in
ΙT
     3190-70-3DP, L-Leucine N-carboxyanhydride, graft polymers with
     butylcarbonyl-protected polyallylamine
                                              24424-99-5DP,
     Di-tert-butyl dicarbonate, reaction products with polyallylamine
       graft polymers with L-leucine N-carboxyanhydride, deprotected
     30551-89-4DP, Polyallylamine, reaction products with di-Bu
     dicarbonate, graft polymers with L-leucine N-carboxyanhydride, deprotected
     870636-97-8P
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP
     (Physical process); SPN (Synthetic preparation); PREP (Preparation); PROC
        (preparation and pH-induced reversible conformational and morphol.
        regulation of polyleucine grafted polyallylamine assembly in
        solution)
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     ANSWER 6 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
L8
ΑN
     2005:547065 CAPLUS
DN
     143:73874
ED
     Entered STN:
                   24 Jun 2005
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ΤI
    Catalytic enzyme-modified textiles for active protection from toxins
IN
    Singh, Alok; Dressick, Walter J.; Lee, Yongwoo
PA
SO
    U.S. Pat. Appl. Publ., 18 pp., Sont.-in-part of U.S. Ser. No. 750,637.
    CODEN: USXXCO
DT
    Patent
LA
    English
IC
     ICM C12N009-16
INCL 435196000
     7-7 (Enzymes)
     Section cross-reference(s): 40
FAN.CNT 2
                               DATE
     PATENT NO.
                        KIND
                                           APPLICATION NO.
                                                                  DATE
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                                           ______
                        A1
                               20050623
                                           US 2004-849621
PΙ
    US 2005136523
                                                                  20040520
                        A1
    US 2005136522
                               20050623
                                           US 2003-750637
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    US 7067294 B2
US 2007014838 A1
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US 2004-849621 A3
                               20060627
                               20070118
                                           US 2006-527650
                                                                  20060919
PRAI US 2003-750637
                               20031223
                              20040520
CLASS
                CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
                _____
US. 2005136523
                ICM
                       C12N009-16
                 INCL
                        435196000
                 IPCI
                       C12N0009-16 [ICM, 7]
                 IPCR
                       A61K0038-43 [I,C*]; A61K0038-46 [I,A]; C12N0009-16
                        [I,C*]; C12N0009-16 [I,A]
                 NCL
                        435/196.000
                 ECLA
                        A62D005/00; A61K038/46
US 2005136522
                 IPCI C12N0011-00 [I,A]
                 IPCR
                        A61K0038-43 [I,C*]; A61K0038-46 [I,A]; C12N0009-16
                        [I,C*]; C12N0009-16 [I,A]
                 NCL
                        435/196.000; 424/094.600
                 ECLA
                        A62D005/00; A61K038/46
US 2007014838
                 IPCI
                       A61K0009-70 [I,A]; B32B0027-04 [I,A]; C12N0011-04
                        [I,A]; C12N0011-00 [I,C*]; C12N0009-16 [I,A]
                 IPCR.
                       A61K0009-70 [I,C]; A61K0009-70 [I,A]; B32B0027-04
                        [I,C]; B32B0027-04 [I,A]; C12N0009-16 [I,C];
                        C12N0009-16 [I,A]; C12N0011-00 [I,C]; C12N0011-04 [I,A]
                        424/443.000; 435/182.000; 435/196.000; 442/123.000
AB
     The present invention provides bioactive catalytic enzyme-modified
     textiles for providing protection from chemical exposure that are stable and
     retain their catalytic activity comprise a cloth substrate, at least one
    polyelectrolyte layer, at least one enzyme layer to degrade the chemical
     agent, and an end-capping layer. The textiles of the present invention
     can be used for reusable clothing that decontaminates itself after
    exposure to toxins and can be worn multiple times or for disposable
    clothing and wipes intended for a single use that decontaminate themselves
     without harming the environment. In a preferred embodiment, the present
     invention takes advantage of superior catalytic activity of enzymes by
    immobilizing them within polyelectrolyte multilayers (PEMs). The
     technique for forming multilayers is simple and effective as
     polyelectrolytes of opposing polarity are alternatively deposited through
     neutralization and overcompensation of their charges. A capping agent
     provides stability to the multilayers, keeps enzymes protected
     in adverse working environments, and attracts the toxic agents to
     facilitate contact with the catalytic sites. The present invention
     provides several advantages over the prior art. It leads to enhanced
     enzyme shelf life under normal storage conditions. It allows
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incorporation of multiple components into multilayers to provide add- on capabilities to the packaged system. It is lightwt., robust, sturdy, disposable, self-decontaminating, and cost-effective. It offers versatility as it can be designed for use on various materials. One example demonstrates the preparation of a polyelectrolyte-enzyme multilayer containing a single layer of the OPH enzyme on a woven fiberglass cloth sample by the dip coating method. catalysis enzyme modified textile toxin decontamination protective clothing Chemicals Decontamination Textiles (catalytic enzyme-modified textiles for active protection from toxins) Enzymes, uses RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES (Uses) (catalytic enzyme-modified textiles for active protection from toxins) Fluoropolymers, uses RL: NUU (Other use, unclassified); USES (Uses) (catalytic enzyme-modified textiles for active protection from toxins) Toxins RL: REM (Removal or disposal); PROC (Process) (catalytic enzyme-modified textiles for active protection from toxins) Glass fiber fabrics RL: TEM (Technical or engineered material use); USES (Uses) (catalytic enzyme-modified textiles for active protection from toxins) Glass fibers, uses RL: TEM (Technical or engineered material use); USES (Uses) (catalytic enzyme-modified textiles for active protection from toxins) Polyamide fibers, uses RL: TEM (Technical or engineered material use); USES (Uses) (catalytic enzyme-modified textiles for active protection from toxins) Rayon, uses RL: TEM (Technical or engineered material use); USES (Uses) (catalytic enzyme-modified textiles for active protection from toxins) Polymers, uses RL: NUU (Other use, unclassified); USES (Uses) (charged, substrate capable of adsorbing; catalytic enzyme-modified textiles for active protection from toxins) Safety devices (chemical protective clothing; catalytic enzyme-modified textiles for active protection from toxins) (chemical protective; catalytic enzyme-modified textiles for active protection from toxins) Amines, uses RL: NUU (Other use, unclassified); USES (Uses) (cotton modified with; catalytic enzyme-modified textiles for active protection from toxins) Textiles Textiles (cotton; catalytic enzyme-modified textiles for active protection from toxins) Coating process

(dip, layers are deposited using; catalytic enzyme-modified textiles

(hexyl, outmost capping layer comprises branched polyethylenimine (BPEI) modified with; catalytic enzyme-modified textiles for active

for active protection from toxins)

for Connie

Functional groups

protection from toxins)

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Polyelectrolytes
IT
        (layer; catalytic enzyme-modified textiles for active protection from .
        toxins)
IΤ
     Antibacterial agents
        (outermost capping layer contains; catalytic enzyme-modified textiles
        for active protection from toxins)
IT
     Acrylic polymers, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (polyelectrolyte layer comprises; catalytic enzyme-modified textiles
        for active protection from toxins)
IT
     Coating process
        (spin, layers are deposited using; catalytic enzyme-modified textiles
        for active protection from toxins)
IΤ
     Coating process
        (spray, layers are deposited using; catalytic enzyme-modified textiles
        for active protection from toxins)
IΤ
     Functional groups
        (substrate comprises a material whose surface chemical modified to
        generate; catalytic enzyme-modified textiles for active protection from
        toxins)
ΙT
     Threads
        (woven into a fabric, substrate is; catalytic enzyme-modified textiles
        for active protection from toxins)
     923-26-2, 1,2-Dihydroxypropyl methacrylate
                                                   1760-24-3,
ΙT
                                                    149305-62-4,
     N-[(3-Trimethoxysilyl)propyl]ethylenediamine
     1,2-Dihydroxypropyl 4-vinylbenzyl ether
     RL: NUU (Other use, unclassified); USES (Uses)
        (branched, capping layer comprises; catalytic enzyme-modified textiles
        for active protection from toxins)
ΙT
     12619-70-4, Cyclodextrin
     RL: NUU (Other use, unclassified); USES (Uses)
        (cotton modified with; catalytic enzyme-modified textiles for active
        protection from toxins)
ΙT
     7585-39-9, β-Cyclodextrin
     RL: NUU (Other use, unclassified); USES (Uses)
        (functionalized polyelectrolyte, polyelectrolyte layer comprises;
        catalytic enzyme-modified textiles for active protection from toxins)
     9032-18-2, Organophosphorous acid anhydrolase
ΙT
     RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
     (Uses)
        (organophosphorous acid anhydrolase; catalytic enzyme-modified textiles
        for active protection from toxins)
     117698-12-1, EC 3.1.8.1
ΙT
     RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
        (organophosphorous hydrolase; catalytic enzyme-modified textiles for
        active protection from toxins)
     74-83-9, Methyl bromide, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (outmost capping layer comprises branched polyethylenimine quaternized
        with; catalytic enzyme-modified textiles for active protection from
        toxins)
ΙT
     9002-84-0, Polytetrafluoroethylene
     RL: NUU (Other use, unclassified); USES (Uses)
        (oxidized, substrate comprises; catalytic enzyme-modified textiles for
        active protection from toxins)
IT
     9047-01-2, Phosphotriesterase
     RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
        (phosphotriesterases (PTE) (EC 3.1.8); catalytic enzyme-modified
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textiles for active protection from toxins)
ΙT
     50851-57-5
     RL: NUU (Other use, unclassified); USES (Uses)
        (polyelectrolyte layer comprises, capping layer comprises; catalytic
        enzyme-modified textiles for active protection from toxins)
     9002-98-6
IT
                 9003-47-8, Polyvinylpyridine
                                                25087-26-7, Polymethacrylic
                                           26062-79-3, Polydiallyl
            25191-25-7, Polyvinyl sulfate
     dimethylammonium chloride
                                 26913-06-4, Poly[imino(1,2-ethanediyl)]
     71550-12-4, Polyallylamine hydrochloride
     RL: NUU (Other use, unclassified); USES (Uses)
        (polyelectrolyte layer comprises; catalytic enzyme-modified textiles
        for active protection from toxins)
L8
     ANSWER 7 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
ΑN
     2004:456842 CAPLUS
DN
     141:217198
     Entered STN: 07 Jun 2004
ED
     X-ray magnetic circular dichroism study of gold nanoparticles
TΤ
     protected by polymer
ΑU
     Yamamoto, Y.; Miura, T.; Teranishi, T.; Suzuki, M.; Kawamura, N.;
     Miyagawa, H.; Nakamura, T.; Kobayashi, K.; Hori, H.
     School of Materials Science, Japan Advanced Institute of Science and
CS
     Technology (JAIST), Tatsunokuchi, Ishikawa, 923-1292, Japan
     Journal of Magnetism and Magnetic Materials (2004), 272-276(Suppl. 1),
SO
     E1183-E1184
     CODEN: JMMMDC; ISSN: 0304-8853
     URL: http://www.sciencedirect.com/science/journal/03048853
PB
   Elsevier Science B.V.
DT
     Journal; (online computer file)
LA
     English .
CC
     77-1 (Magnetic Phenomena)
     Section cross-reference(s): 36, 56, 66, 73
AB
     Bulk Au metal is chemical stable and has diamagnetic properties. However,
     these properties might be different in nano-sized region. This article
     reports that the direct observation of spontaneous spin polarization of Au
     nanoparticles employing x-ray MCD (XMCD). Magnetization estimated by XMCD
     shows a good agreement with the results obtained by d.c. magnetization.
     This result evidences the intrinsic spin polarization in nano-sized
     material.
ST
     gold nanoparticle XMCD XAS magnetization polymer host.
ΤT
     Nanoparticles
     Spin polarization
     X-ray magnetic circular dichroism
        (XMCD study of gold nanoparticles protected by polymer)
ΙT
     Magnetization
     X-ray spectra
        (of gold nanoparticles protected by polymer)
ΙT
     71550-12-4, Polyallylamine hydrochloride
     RL: NUU (Other use, unclassified); USES (Uses)
      (XMCD study of gold nanoparticles protected by polymer)
     7440-57-5, Gold, properties
     RL: PRP (Properties)

    (XMCD study of gold nanoparticles protected by polymer)

RE.CNT 5
              THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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ANSWER 8 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
L8
     2004:236338 CAPLUS
AN
DN
     140:387786
     Entered STN: 23 Mar 2004
ΕD
ΤI
     A simple protocol for the modular assembly of "millipede" artificial
ΑU
     Motherwell, William B.; Atkinson, Catherine E.; Aliev, Abil E.; Wong,
     Stephanie Y. F.; Warrington, Brian H.
CS
     Department of Chemistry, University College London, London, WC1H 0AJ, UK
     Angewandte Chemie, International Edition (2004), 43(10), 1225-1228
SO
     CODEN: ACIEF5; ISSN: 1433-7851
PB
     Wiley-VCH Verlag GmbH & Co. KGaA
DT
     Journal
     English
LΑ
CC
     7-8 (Enzymes)
     Section cross-reference(s): 35
OS
     CASREACT 140:387786
AΒ
     A modular "step-by-step" strategy, which combines both design and
     selection elements for the attachment of sep. considered receptor sites
     and catalytically active groups to a flexible polymer backbone, can be
     used to construct an artificial enzyme. The method outlined is clearly
     amenable to a parallel synthesis approach for optimization and can also
     provide, at every step along the way, detailed information about the
     relative importance and cooperativity of the groups contributing to the
     mol. environment of active sites within enzymes.
     artificial enzyme modular assembly transition state functional group
ST
     polymer; esterase artificial modular assembly transition state functional
     group polymer
İT
     Molecular association
        (esterase transition state analog interaction with dipeptides in
        relation to assembly of artificial enzymes)
     Enzymes, biological studies
TΤ
     RL: BSU (Biological study, unclassified); CAT (Catalyst use); SPN
     (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (synthetic; modular step-by-step protocol for assembly of artificial
        enzymes based on transition state analog-related receptor sites and
        catalytically active residues on flexible polymer)
                           687-69-4
                                                  16709-12-9
ΙT
     305-84-0
                658-79-7
                                      7093-70-1
                                                                17136-25-3
     22467-93-2
                  67726-09-4
     RL: BSU (Biological study, unclassified); PEP (Physical, engineering or
     chemical process); PYP (Physical process); BIOL (Biological study); PROC
     (Process)
        (esterase transition state analog interaction with dipeptides in
        relation to assembly of artificial enzymes)
     71550-12-4DP, reaction products with protected lysine and
                       78081-87-5DP, reaction products with
     arginylarginine
     polyallylamine and protected arginylarginine
     688044-44-2DP, reaction products with polyallylamine and
     protected lysine
     RL: BSU (Biological study, unclassified); CAT (Catalyst use); SPN
     (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (modular step-by-step protocol for assembly of artificial esterases
        based on transition state analog-related receptor sites and
```

catalytically active residues on flexible polymer)

(modular step-by-step protocol for assembly of artificial esterases

688044-44-2

71550-12-4, Polyallylamine hydrochloride

RL: RCT (Reactant); RACT (Reactant or reagent)

IT

```
based on transition state analog-related receptor sites and
        catalytically active residues on flexible polymer)
IT
     588708-82-1
     RL: BSU (Biological study, unclassified); RCT (Reactant); BIOL (Biological
     study); RACT (Reactant or reagent)
         (substrate ester; modular step-by-step protocol for assembly of
        artificial esterases based on transition state analog-related receptor
        sites and catalytically active residues on flexible polymer)
IT
     588708-83-2
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); BIOL (Biological study); USES (Uses)
         (transition state analog; modular step-by-step protocol for assembly of
        artificial esterases based on transition state analog-related receptor
        sites and catalytically active residues on flexible polymer)
RE.CNT
              THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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     ANSWER 9 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
L8
     2003:747019 CAPLUS
ΑN
DN
     139:263806
ED
     Entered STN:
                    24 Sep 2003
     Formation of Silver Nanoparticles under Structured Amino Groups in
TI
     Pseudo-dendritic Poly(allylamine) Derivatives
ΑU
     Kuo, Ping-Lin; Chen, Wei-Fu
     Department of Chemical Engineering, National Cheng Kung University,
CS
     Tainan, 70101, Taiwan
     Journal of Physical Chemistry B (2003), 107(41), 11267-11272
SO
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CODEN: JPCBFK; ISSN: 1520-6106
PB
     American Chemical Society
DT
     Journal
LA
     English
CC
     56-1 (Nonferrous Metals and Alloys)
     Section cross-reference(s): 38
AB
     The syntheses of silver nanoparticles stabilized by poly(allylamine) (PAA)
     and by polyethyleneiminated poly(allylamine) (PAA(EI)n (n = 2, 5.8)) are
     reported. The architectural effects in particle on the nanoparticle size,
     size distribution, and agglomeration behavior are determined from the UV-vis
     plasmon absorption band and transmission electron microscopic (TEM)
     analyses. The data show that PAA(EI)n display better stabilizing effects
     than PAA to prevent silver particles from agglomeration. Different
     phenomena of the polymer-protected nanoparticles at various
     silver ion concns. are observed and are explained in terms of a mechanism of
     structure-dependent stabilization.
ST
     silver nanoparticle polyallylamine deriv
   Nanoparticles
         (formation of Silver Nanoparticles under Structured Amino Groups in
        Pseudo-dendritic Poly(allylamine) Derivs.)
IT
     7440-22-4P, Silver, preparation
     RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or
     engineered material use); PREP (Preparation); USES (Uses)
         (formation of Silver Nanoparticles under Structured Amino Groups in
        Pseudo-dendritic Poly(allylamine) Derivs.)
IΤ
     30551-89-4, Polyallylamine
     RL: CPS (Chemical process); NUU (Other use, unclassified); PEP (Physical,
     engineering or chemical process); PROC (Process); USES (Uses)
         (polyethyleneiminated; formation of Silver Nanoparticles under
        Structured Amino Groups in Pseudo-dendritic Poly(allylamine) Derivs.)
RE.CNT
              THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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L8
     ANSWER 10 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
ΑN
     2003:551891 CAPLUS
DN
     139:231525
ED
     Entered STN: 20 Jul 2003
TΙ
     Preparation of polyelectrolyte-layered assemblies containing cyclodextrin
     and their binding properties
ΑU
     Sato, Katsuhiko; Suzuki, Iwao; Anzai, Jun-ichi
     Graduate School of Pharmaceutical Sciences, Tohoku University, Sendai,
CS
     908-8578, Japan
     Langmuir (2003), 19(18), 7406-7412
SO
     CODEN: LANGD5; ISSN: 0743-7463
PB
     American Chemical Society
DT
     Journal
LA
     English
     38-2. (Plastics Fabrication and Uses)
CC
     Section cross-reference(s): 41
AB
     Polyelectrolyte-layered films containing \alpha- and \beta-cyclodextrin were
     prepared by a layer-by-layer deposition of sulfonated \alpha- or
     \beta-cyclodextrin (s-\alpha-CyD or s-\beta-CyD) and cationic
     poly(allylamine hydrochloride) (PAH) on a solid surface. The deposition
     of the films from water solns. (no salt added) afforded thin films in
     which s\text{-}\alpha\text{-}\text{CyD} and s\text{-}\beta\text{-}\text{CyD} formed roughly a monomol. layer upon
     each deposition, whereas the films were thicker when a high-ionic-strength-
     buffer was employed as a bathing solution. The PAH-s-\alpha-CyD film bound
     methyl orange (MO) by forming an inclusion complex in the film, whereas MO
     did not form the complex with s-\beta-CyD in the PAH-s-\beta-CyD film
     but formed aggregates on the surface of the film. MO was not desorbed
     from the films upon rinsing in pure water or even in a HCl solution (pH 2.0),
     whereas MO was extracted from the film into a NaOH solution (pH 12.0). MO
     complexed with s-\alpha-CyD in the film was protected from
     protonation in HCl solution (pH 2.0). The PAH-s-\alpha-CyD film showed a
     binding selectivity to azoarom. dyes, depending on the size and shape of
     the dyes.
ST
     cyclodextrin polyallylamine polyelectrolyte layered sensor
     assembly prepn dye binding
ΙT
     Azo dyes
         (aromatic; polyelectrolyte-layered assemblies containing cyclodextrins and
       their dye-binding properties for use as chemical sensors)
IT
         (layered assemblies; polyelectrolyte-layered assemblies containing
        cyclodextrins and their dye-binding properties for use as chemical
        sensors)
IT
     Monolayers
     Polyelectrolytes
         (polyelectrolyte-layered assemblies containing cyclodextrins and their
        dye-binding properties for use as chemical sensors)
     71550-12-4, Poly(allylamine hydrochloride
IT
     RL: NUU (Other use, unclassified); USES (Uses)
         (cationic; polyelectrolyte-layered assemblies containing cyclodextrins and
        their dye-binding properties for use as chemical sensors)
     7585-39-9D, \beta-Cyclodextrin, sulfonated
                                                10016-20-3D,
ΙT
     \alpha-Cyclodextrin, sulfonated
     RL: NUU (Other use, unclassified); USES (Uses)
         (polyelectrolyte-layered assemblies containing cyclodextrins and their
        dye-binding properties for use as chemical sensors)
     547-58-0, Methyl orange
ΙT
     RL: NUU (Other use, unclassified); PRP (Properties); USES (Uses)
        (polyelectrolyte-layered assemblies containing cyclodextrins and their
        dye-binding properties for use as chemical sensors)
              THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS RECORD
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L8
     ANSWER 11 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
     2003:243023 CAPLUS
ΑN
     139:96917
DN
ED
     Entered STN:
                   30 Mar 2003
TΤ
     Structural regulation of a peptide-conjugated graft copolymer: A simple
     model for amyloid formation
ΑU
     Koga, Tomoyuki; Taguchi, Kazuhiro; Kobuke, Yoshiaki; Kinoshita, Takatoshi;
     Higuchi, Masahiro
CS
     Nanoarchitectonics Research Center, National Institute of Advanced
     Industrial Science and Technology and CREST (Japan Science and
     Technology), Ibaraki, 305-8565, Japan
```

Chemistry--A European Journal (2003), 9(5), 1146-1156

CODEN: CEUJED; ISSN: 0947-6539

Wiley-VCH Verlag GmbH & Co. KGaA

Journal

SO

PΒ

DΤ

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11/245136:
LA
     English
CC
     6-3 (General Biochemistry)
OS
     CASREACT 139:96917
AΒ
     The self-assembly of peptides and proteins into \beta-sheet-rich
     high-order structures has attracted much attention as a result of the
     characteristic nanostructure of these assemblies and because of their
     association with neurodegenerative diseases. Here we report the structural
     and conformational properties of a peptide-conjugated graft copolymer,
     poly(\gamma-methyl-L-glutamate) grafted polyallylamine (1) in a
     water-2,2,2-trifluoroethanol solution as a simple model for amyloid
     formation. Atomic force microscopy revealed that the globular peptide 1
     self-assembles into nonbranching fibrils that are about 4 nm in height
     under certain conditions. These fibrils are rich in \beta-sheets and,
     similar to authentic amyloid fibrils, bind the amyloidophilic dye Congo
           The secondary and quaternary structures of the peptide 1 can be
     controlled by manipulating the pH, solution composition, and salt
concentration; this
     indicates that the three-dimensional packing arrangement of peptide chains
     is the key factor for such fibril formation. Furthermore, the addition of
     carboxylic acid-terminated poly(ethylene glycol), which interacts with
     both of amino groups of 1 and hydrophobic PMLG chains, was found to
     obviously inhibit the \alpha\text{-to-}\beta structural transition for
     non-assembled peptide 1 and to partially cause a \beta\text{-to-}\alpha
     structural transition against the 1-assembly in the \beta\text{--sheet} form.
     These findings demonstrate that the amyloid fibril formation is not
     restricted to specific protein sequences but rather is a generic property
     of peptides. The ability to control the assembled structure of the
     peptide should provide useful information not only for understanding the
     amyloid fibril formation, but also for developing novel peptide-based
     material with well-defined nanostructures.
ST
     peptide conjugated graft copolymer amyloid fibril model
ΙT
     Fibril
        (amyloid-like; peptide-conjugated graft copolymer as a simple model for
        amyloid fibril formation)
IT
     Organelle
       . (fibril, amyloid-like; peptide-conjugated graft copolymer as a simple
        model for amyloid fibril formation)
IT
     Polymers, biological studies
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (graft; peptide-conjugated graft copolymer as a simple model for
        amyloid fibril formation)
ΙT
     Self-assembly
        (into amyloid-like fibrils; peptide-conjugated graft copolymer as a
        simple model for amyloid fibril formation)
ΙT
     Conformation
     β-Sheet
        (peptide-conjugated graft copolymer as a simple model for amyloid
        fibril formation)
ΙT
     Amyloid
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (peptide-conjugated graft copolymer as a simple model for amyloid
        fibril formation)
ΙT
     36877-69-7DP, reaction products with allylamine-Me glutamate graft
     copolymer/nitrobenzofurazan derivs.
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (Rhodamine B-containing peptide-conjugated graft copolymer;
```

peptide-conjugated graft copolymer as a simple model for amyloid fibril

formation)

```
29270-56-2DP, reaction products with allylamine-Me glutamate graft
     copolymer
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (nitrobenzofurazan-containing peptide-conjugated graft copolymer;
        peptide-conjugated graft copolymer as a simple model for amyloid fibril
        formation)
     1663-47-4, \gamma-Methyl-L-glutamate-N-carboxy anhydride
ΙT
                                                            24424-99-5,
     Di-tert-butyl dicarbonate
                                 29270-56-2, 4-Fluoro-7-nitrobenzofurazan
     30551-89-4, Polyallylamine
                                  36877-69-7, Rhodamine B
     isothiocyanate
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (peptide-conjugated graft copolymer as a simple model for amyloid
        fibril formation)
ΙT
     431047-85-7DP, BOC-protected, benzofurazan derivs., reaction
     products with Rhodamine B isothiocyanate
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (peptide-conjugated graft copolymer; peptide-conjugated graft copolymer
        as a simple model for amyloid fibril formation)
RE.CNT
              THERE ARE 61 CITED REFERENCES AVAILABLE FOR THIS RECORD
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- L8 ANSWER 12 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
- AN 2002:885538 CAPLUS
- DN 139:85620
- ED Entered STN: 22 Nov 2002
- TI Spontaneous formation of fibrillar β -sheet assemblies from peptide-grafted polyamine; effect of complexation with poly(ethylene qlycol) derivatives
- AU Koqa, Tomoyuki; Taguchi, Kazuhiro; Higuchi, Masahiro; Kinoshita, Takatoshi
- CS Nanoarchitectonics Research Center, National Institute of Advanced Industrial Science and Technology and CREST (Japan Science and Technology), Tsukuba, Ibaraki, 305-8565, Japan
- SO Transactions of the Materials Research Society of Japan (2002), 27(3), 493-496
 CODEN: TMRJE3; ISSN: 1382-3469
- PB Materials Research Society of Japan
- DT Journal
- LA English
- CC 34-3 (Amino Acids, Peptides, and Proteins) Section cross-reference(s): 22, 35
- OS CASREACT 139:85620
- Amphiphilic graft copolymer, polyallylamine bearing poly(γ -methyl-L-glutamate) graft chains (PAAgPMLG), was prepared The conformation of PAAgPMLG was changed spontaneously from α -helix to β -sheet only when the amino groups of PAA units were protonated (pH < 8). In this condition, PAAgPMLG formed amyloid-like fibrils with regular quaternary structure. On the other hand, such fibril formation with α -to- β structural transition was obviously inhibited in the presence of carboxylic acid-terminated poly(ethylene glycol), which prevents appropriate folding of peptide graft-chain through the complexation with amino groups of PAAgPMLG. These findings demonstrate that the amyloid fibril formation seems to occur if peptide chains are folded appropriately even at the simple synthetic peptide, which have no specific protein sequences.
- ST fibril beta sheet polyglutamate grafted polyallylamine effect

```
PEG deriv
 ΙT
      Polyoxyalkylenes, usės
      RL: MOA (Modifier or additive use); USES (Uses)
          (amine-terminated; spontaneous formation of fibrillar \beta-sheet
         assemblies from polyglutamate-grafted polyallylamine and
          inhibition in the presence of PEG derivs.)
       Polyoxyalkylenes, uses
 ΙT
      RL: MOA (Modifier or additive use); USES (Uses)
          (carboxy-terminated; spontaneous formation of fibrillar \beta-sheet
          assemblies from polyglutamate-grafted polyallylamine and
          inhibition in the presence of PEG derivs.)
 ΙT
      Conformation
      Fibril
      B-Sheet
          (spontaneous formation of fibrillar \beta-sheet assemblies from
         polyglutamate-grafted polyallylamine and inhibition in the
         presence of PEG derivs.)
 ΙT
      Amyloid
      RL: BSU (Biological study, unclassified); BIOL (Biological study)
          (spontaneous formation of fibrillar \beta-sheet assemblies from
         polyglutamate-grafted polyallylamine and inhibition in the
          presence of PEG derivs.)
 ΙT
       25322-68-3D, amine-terminated
                                       25322-68-3D, carboxy-terminated
      RL: MOA (Modifier or additive use); USES (Uses)
          (spontaneous formation of fibrillar \beta-sheet assemblies from
         polyglutamate-grafted polyallylamine and inhibition in the
         presence of PEG derivs.)
· IT
       431047-85-7P
      RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
          (spontaneous formation of fibrillar \beta-sheet assemblies from
          polyglutamate-grafted polyallylamine and inhibition in the
          presence of PEG derivs.)
 ΙT
       1663-47-4
                   30551-89-4, Polyallylamine
       RL: RCT (Reactant); RACT (Reactant or reagent)
          (spontaneous formation of fibrillar \beta-sheet assemblies from
          polyglutamate-grafted polyallylamine and inhibition in the
          presence of PEG derivs.)
 ΙT
       30551-89-4DP, BOC-protected
       RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
       (Reactant or reagent)
          (spontaneous formation of fibrillar \beta-sheet assemblies from
          polyglutamate-grafted polyallylamine and inhibition in the
          presence of PEG derivs.)
 RE.CNT
          22
                THERE ARE 22 CITED REFERENCES AVAILABLE FOR THIS RECORD
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(17) Sasaki, T; J Am Chem Soc 1989, V111, P380 CAPLUS
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(22) Yamada, N; J Am Chem Soc 1998, V120, P12192 CAPLUS
     ANSWER 13 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
L8
ΑN
     2002:106264 CAPLUS
DN
     136:295311
ED
     Entered STN: 10 Feb 2002
TI.
     Layer-by-layer growth of polymer/nanoparticle films containing monolayer-
     protected gold clusters
ΑU
     Hicks, Jocelyn F.; Young, Seok-Shon; Murray, Royce W.
CS
     Kenan Laboratories of Chemistry, University of North Carolina, Chapel
     Hill, NC, 27599-3290, USA
SO
     Langmuir (2002), 18(6), 2288-2294
     CODEN: LANGD5; ISSN: 0743-7463
PΒ
     American Chemical Society.
DT
     Journal
LA
     English
     36-5 (Physical Properties of Synthetic High Polymers)
CC
     Section cross-reference(s): 76
AΒ
     Multilayer films of nanoparticles were grown in a systematic and
     controlled manner layer-by-layer by alternating exposures of suitably
     functionalized substrates (glass, Au) to either poly(allylamine) and
     carboxylic acid-functionalized nanoparticles or to poly(styrene sulfonate)
     and arylamine-functionalized nanoparticles. Electrostatic interactions
     comprise the dominant film growth factors. The rate of multilayer film
     growth depends on the polymer solution pH and other details of the solution
     exposures. Growth was followed by spectrophotometry of the Au
     nanoparticle cores, voltammetry of the Au core double layer charging, and
     film mass (quartz crystal microbalance). The first example is reported of
     quantized double layer charging of the Au cores in a layer-by-layer film
     that is composed of monolayer-protected clusters and a
     polyelectrolyte.
ST.
     polyallylamine gold nanoparticle layer deposition electron
     transport double layer; polystyrene sulfonate gold nanoparticle multilayer
     electron transport double layer
ΙT
     Adsorbed monolayers
     Electric capacitance
     Electric double layer
     Electron transport
     Laminated plastic films
     Polyelectrolytes
        (layer-by-layer growth of polymer/nanoparticle films containing monolayer-
        protected gold clusters)
ΙT
     1193-02-8, 4-Aminothiophenol
                                     71310-21-9, 11-Mercaptoundecanoic acid
     RL: NUU (Other use, unclassified); USES (Uses)
        (layer-by-layer growth of polymer/nanoparticle films containing monolayer-
        protected gold clusters)
     25704-18-1, Poly(sodium 4-styrene sulfonate)
                                                      71550-12-4,
IT
     Polyallylamine hydrochloride
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP
     (Physical process); PROC (Process)
         (layer-by-layer growth of polymer/nanoparticle films containing monolayer-
        protected gold clusters)
ΙT
     7440-57-5, Gold, processes
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
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process); PROC (Process)
         (layer-by-layer growth of polymer/nanoparticle films containing monolayer-
         protected gold clusters)
RE.CNT
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     ANSWER 14 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
F8 ·
AN ·
     2002:97735 CAPLUS
DN
     136:402005
ED
     Entered STN: 06 Feb 2002
TI
     pH-Regulated formation of amyloid-like β-sheet assemblies from
     polyglutamate grafted polyallylamine
     Koga, Tomoyuki; Taguchi, Kazuhiro; Kinoshita, Takatoshi; Higuchi, Masahiro
ΑU
CS
     Nanoarchitectonics Research Center, National Institute of Advanced
     Industrial Science and Technology and CREST, Japan Science and Technology,
     Ibaraki, Tsukuba, 305-8565, Japan
SO
     Chemical Communications (Cambridge, United Kingdom) (2002), (3), 242-243
     CODEN: CHCOFS; ISSN: 1359-7345
PΒ
     Royal Society of Chemistry
DT
     Journal
     English
LA
CC
     34-3 (Amino Acids, Peptides, and Proteins)
     Section cross-reference(s): 22, 35
AB
     A novel artificial protein with simple primary structure,
     poly(\gamma-methyl-L-glutamate)-grafted polyallylamine, has
     been prepared and the resultant peptide has shown a unique property of
     pH-regulated conformation and morphol.
ST
     pH amyloid like beta sheet polyglutamate grafted polyallylamine
ΙT
     Conformation
     рΗ
     β-Sheet
        (pH-regulated formation of amyloid-like \beta-sheet assemblies from
        polyglutamate grafted polyallylamine)
IT
     431047-85-7P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (pH-regulated formation of amyloid-like \beta-sheet assemblies from
        polyglutamate grafted polyallylamine)
     1663-47-4
IT
                 30551-89-4, Polyallylamine
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (pH-regulated formation of amyloid-like \beta-sheet assemblies from
        polyglutamate grafted polyallylamine)
ΙT
     30551-89-4DP, BOC-protected
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (pH-regulated formation of amyloid-like \beta-sheet assemblies from
        polyglutamate grafted polyallylamine)
RE.CNT
              THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD
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(13) Miyazawa, T; J Am Chem Soc 1961, V83, P712 CAPLUS
(14) Mutter, M; Angew Chem, Int Ed Engl 1989, V28, P535
(15) Ramirez-Aguilar, K; Langmuir 1998, V14, P2562 CAPLUS
(16) Rochet, J; Curr Opin Struc Biol 2000, V10, P60 CAPLUS
(17) Sasaki, T; J Am Chem Soc 1989, V111, P380 CAPLUS (18) Sipe, J; Crit Rev Clin Lab Sci 1994, V31, P325 CAPLUS
(19) Takahashi, Y; Chem Eur J 1998, V12, P2475 (20) Takahashi, Y; Structure 2000, V8(9), P915 CAPLUS
(21) Wallimann, P; Angew Chem, Int Ed 1999, V38, P1290 CAPLUS
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     ANSWER 15 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
\Gamma8
AN
     2000:738734 CAPLUS
DN
     133:310972
ED
     Entered STN: 19 Oct 2000
TΙ
     Metal articles protected with crosslinked polyamine coating
     Keller, Harald; Jaworek, Thomas; Schliephake, Volker; Schroeder, Juergen;
TN
     Sutoris, Heinz Friedrich; Wagenblast, Gerhard
PΑ
     BASF A.-G., Germany
SO
     Ger. Offen., 6 pp.
     CODEN: GWXXBX
DT
   Patent
LA
     German
IC
     ICM C09D179-02
     ICS C09D131-02; C09D125-04; B05D007-16; B65D025-14; B65D090-04;
          B01J019-02
     42-10 (Coatings, Inks, and Related Products)
CC
FAN.CNT 1
     PATENT NO.
                                              APPLICATION NO.
                                                                      DATE
                          KIND
                                DATE
                                              ------
     _____
                          ____
                                                                      _____
                                 _____
     DE 19915116
                                              DE 1999-19915116
                                                                      19990401
                          A1
                                  20001019
PΙ
PRAI DE 1999-19915116
                                 19990401
CLASS
 PATENT NO.
                  CLASS PATENT FAMILY CLASSIFICATION CODES
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DE 19915116
                  ICM
                         C09D179-02
                         C09D131-02; C09D125-04; B05D007-16; B65D025-14;
                  ICS
                         B65D090-04; B01J019-02
                         C09D0179-02 [ICM,7]; C09D0179-00 [ICM,7,C*];
                  IPCI
                         C09D0131-02 [ICS, 7]; C09D0131-00 [ICS, 7, C*];
                         C09D0125-04 [ICS,7]; C09D0125-00 [ICS,7,C*];
                         B05D0007-16 [ICS,7]; B65D0025-14 [ICS,7]; B65D0090-04
                         [ICS, 7]; B01J0019-02 [ICS, 7]
                  IPCR
                         B01J0019-02 [I,C*]; B01J0019-02 [I,A]; B05D0003-10
                         [I,C*]; B05D0003-10 [I,A]; B05D0007-16 [I,C*];
                         B05D0007-16 [I,A]; C09D0179-00 [I,C*]; C09D0179-02
                  ECLA
                         B01J019/02; B05D003/10L3; B05D007/16; C09D179/02
AB
     Metal articles coated with crosslinked polyamine coatings are claimed.
     The coatings protect the inner walls of metal containers and reactors
     against unwanted accumulation of polymers. Thus, a stainless steel net
     was coated by immersing in aqueous polyethyleneimine solution followed by
     immersing in aqueous glutaraldehyde solution and drying at ambient temperature
ST
     polyamine crosslinked coating metal surface protection; coating
     polyethyleneimine metal surface protection; glutaraldehyde crosslinker
     polyethyleneimine coating metal surface protection
```

```
ΙT
    Coating materials
        (metal articles protected with crosslinked polyamine coating)
ΙT
    Polvamines
    RL: TEM (Technical or engineered material use); USES (Uses)
        (polyalkylene-; metal articles protected with crosslinked
       polyamine coating)
ΙT
    25086-42-4, Poly(4-aminostyrene) 25104-18-1, Polylysine
                                                                30551-89-4,
    Polyallylamine 38000-06-5, Polylysine 75133-03-8,
    Aziridine-Glutaraldehyde copolymer 111616-54-7, Ethenamine-N-
    Vinylformamide copolymer
    RL: TEM (Technical or engineered material use); USES (Uses)
        (crosslinked; metal articles protected with crosslinked
       polyamine coating)
    12597-68-1, Stainless steel, processes
IT
    RL: PEP (Physical, engineering or chemical process); PROC (Process)
        (mesh; metal articles protected with crosslinked polyamine
       coating)
    7439-89-6, Iron, miscellaneous 7439-98-7, Molybdenum, miscellaneous
ΙT
    7440-02-0, Nickel, miscellaneous 7440-32-6, Titanium, miscellaneous
    7440-47-3, Chromium, miscellaneous 7440-48-4, Cobalt, miscellaneous
    7440-62-2, Vanadium, miscellaneous
    RL: MSC (Miscellaneous)
        (metal articles protected with crosslinked polyamine coating)
    26336-38-9, Polyvinylamine 75133-03-8D, Aziridine-Glutaraldehyde
IT
    copolymer, reduced
    RL: TEM (Technical or engineered material use); USES (Uses)
        (metal articles protected with crosslinked polyamine coating)
    ANSWER 16 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
L8
ΑN
    1998:650992 CAPLUS
DN
    129:332238
ED
    Entered STN: 14 Oct 1998
    Treatment of porous articles and microbiocidal and insect-repellent
ΤI
    compositions containing boron compounds for them
    Aoki, Hiroshi; Tanaka, Kazumi; Echigo, Takashi
ΙN
    Showa Denko K. K., Japan
PA
     Jpn. Kokai Tokkyo Koho, 27 pp.
SO
    CODEN: JKXXAF
DΤ
    Patent
LA
     Japanese
IC
     ICM C08F002-00
     ICS B05D005-00; B05D007-00; B27K003-02; B27K003-34; B27K003-52;
         · C08G073-00
CC
     42-10 (Coatings, Inks, and Related Products)
     Section cross-reference(s): 5, 16, 43
FAN.CNT 1
                                           APPLICATION NO.
     PATENT NO.
                        KIND
                               DATE
                                                                 DATE
                                           -----
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                        ____
                                           JP 1997-75267
                                                                 19970327
    JP 10265508
                         Α
                               19981006
PRAI JP 1997-75267
                               19970327
CLASS
 PATENT NO.
                CLASS · PATENT FAMILY CLASSIFICATION CODES
                ____
                       _____
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 JP 10265508
                ICM
                       C08F002-00
                       B05D005-00; B05D007-00; B27K003-02; B27K003-34;
                ICS
                       B27K003-52; C08G073-00
                       C08F0002-00 [ICM, 6]; B05D0005-00 [ICS, 6]; B05D0007-00
                IPCI
                        [ICS, 6]; B27K0003-02 [ICS, 6]; B27K0003-34 [ICS, 6];
                       B27K0003-52 [ICS,6]; C08G0073-00 [ICS,6]
               · IPCR
                       B05D0005-00 [I,A]; B05D0005-00 [I,C*]; B05D0007-00
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[I,A]; B05D0007-00 [I,C*]; B27K0003-02 [I,A];
                         B27K0003-02 [I,C*]; B27K0003-34 [I,A]; B27K0003-34
                         [I,C*]; B27K0003-52 [I,A]; B27K0003-52 [I,C*];
                         C08F0002-00 [I,A]; C08F0002-00 [I,C*]; C08G0073-00
                         [I,A]; C08G0073-00 [I,C*]
      Porous articles are immersed in and/or coated with organic compds. having
 AB
      CH2:CH, CH:CH, CH2:CMe, CH2:N, CH:N, NH:CH, NH:N and/or N:N (A), B-containing
      compds. (B), and polymerization catalysts (C) and polymerized Compns. for the
      articles comprise A, B, and C. Thus, 1% saponified polymer prepared from
      acetal protected 4-allyl-1,2-dihydroxybenzene and vinyl acetate,
      0.6% boric acid, 2% ligninsulfonic acid, 30 ppm polyphenol oxidase was
      blended to give a solution, in which a chip of wood was immersed. After the
      solution in the chip was oxidized, polymerized, and mixed with H2O, 8% boric
 acid
      flowed out from the chip.
      vinyl alc polymer coating porous article; boron blend polymer coating
 ST
      wood; boric acid polymer coating bactericidal fungicidal; rust inhibiting
      boric acid vinyl coating; insect repellent coating vinyl polymer
 TΤ
      Coating materials
         (anticorrosive; treatment of porous articles with microbiocidal and
        insect-repellent compns. containing boron compds.)
· IT
      Coating materials
         (bactericidal; treatment of porous articles with microbiocidal and
         insect-repellent compns. containing boron compds.)
 IT
      Insect repellents
         (coatings; treatment of porous articles with microbiocidal and
         insect-repellent compns. containing boron compds.)
 IT
      Coating materials
      Coating materials
         (fungicidal; treatment of porous articles with microbiocidal and
         insect-repellent compns. containing boron compds.)
 IT
      Enzymes, uses
      RL: CAT (Catalyst use); USES (Uses)
         (metallo-, polymerization catalysts; treatment of porous articles with
         microbiocidal and insect-repellent compns. containing boron compds.)
 IT
      Transition metals, uses
      RL: CAT (Catalyst use); USES (Uses)
         (polymerization catalysts; treatment of porous articles with microbiocidal
 and
         insect-repellent compns. containing boron compds.)
 IT
      Polymerization catalysts
      Wood
         (treatment of porous articles with microbiocidal and insect-repellent
         compns. containing boron compds.)
      7439-89-6, Iron, uses 7439-92-1, Lead, uses 7439-95-4, Magnesium, uses
 ΙT
      7439-98-7, Molybdenum, uses
                                    7440-02-0, Nickel, uses
                                                              7440-05-3,
      Palladium, uses
                        7440-22-4, Silver, uses
                                                  7440-31-5, Tin, uses
      7440-32-6, Titanium, uses
                                  7440-36-0, Antimony, uses
                                                              7440 - 38 - 2,
                                                  7440-48-4, Cobalt, uses
      Arsenic, uses
                      7440-47-3, Chromium, uses
      7440-50-8, Copper, uses
                                7440-67-7, Zirconium, uses
      RL: BAC (Biological activity or effector, except adverse); BSU (Biological
      study, unclassified); BUU (Biological use, unclassified); MOA (Modifier or
      additive use); BIOL (Biological study); USES (Uses)
         (additives; treatment of porous articles with microbiocidal and
        insect-repellent compns. containing boron compds.)
                                                                    7440-48-4D,
 IT
      87-69-4D, cobalt complex, uses 7429-90-5, Aluminum, uses
      Cobalt, L-tartaric acid complex, uses
                                              7440-66-6, Zinc, uses
                                                                       9002-10-2,
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9003-99-0, Peroxidase

80619-01-8, Bilirubin oxidase

9029-44-1, Ascorbate oxidase

Catechol oxidase

80498-15-3, Laccase

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RL: CAT (Catalyst use); USES (Uses)
        (polymerization catalyst; treatment of porous articles with microbiocidal
and
        insect-repellent compns. containing boron compds.)
ΙT
     87-66-1DP, Pyrogallol, protected, polymer with vinyl acetate and
     ligninsulfonic acid, saponified 93-15-2DP, 4-Allyl-1,2-dimethoxybenzene,
     protected, polymer with vinyl acetate and ligninsulfonic acid,
     saponified 108-05-4DP, Acetic acid ethenyl ester, polymer with dihydroxy
     compds. and ligninsulfonic acid, saponified, uses
                                                         8062-15-5DP,
     Ligninsulfonic acid, polymer with vinyl acetate and dihydroxy compds.,
     saponified
                  58169-20-3DP, protected, polymer with vinyl acetate
     and ligninsulfonic acid, saponified 214957-15-0DP, 4-Allyl-1,2-
     dimethoxybenzene-lignosulfonic acid-vinyl acetate copolymer, saponified
     214957-16-1P, 3,4-Dihydroxybenzaldehyde-ligninsulfonic
     acid-polyethyleneimine copolymer
                                        214957-17-2P, 3,4-Dihydroxybenzaldehyde-
     ligninsulfonic acid-polyallylamine copolymer
                                                    214957-18-3P ·
     214957-19-4P
                    214957-20-7P
                                  215231-98-4P .215232-45-4P
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); IMF (Industrial
     manufacture); POF (Polymer in formulation); PRP (Properties); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (treatment of porous articles with microbiocidal and insect-repellent
        compns. containing boron compds.)
IΤ
     118-79-6, 2,4,6-Tribromophenol
                                     154-23-4, (+)-Catechin
                                                               499-44-5,
                  7173-51-5, Didecyldimethylammonium chloride
     Hinokitiol
                                                                7585-20-8,
     Zirconium acetate. 7758-98-7, Copper sulfate, uses
                                                           7785-70-8
     10025-91-9, Antimony trichloride 10043-35-3, Boric acid (H3BO3), uses
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); MOA (Modifier or
     additive use); BIOL (Biological study); USES (Uses)
        (treatment of porous articles with microbiocidal and insect-repellent
        compns. containing boron compds.)
ΙT
     14475-11-7, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (treatment of porous articles with microbiocidal and insect-repellent
        compns. containing boron compds.)
L8
     ANSWER 17 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
ΑN
     1993:125076 CAPLUS
DN
     118:125076
ED
     Entered STN: 30 Mar 1993
TI
     Preparation of peptide derivatives and their application as antitumor
IN
     Kitaquchi, Hiroshi; Komazawa, Hiroyuki; Kojima, Masayoshi; Mori, Hideto;
     Nishikawa, Naoyuki; Satoh, Hideaki; Orikasa, Atsushi; Ono, Mitsunori;
     Azuma, Ichiro; Saiki, Ikuo
PΑ
     Fuji Photo Film Co., Ltd., Japan
SO
     Eur. Pat. Appl., 69 pp.
     CODEN: EPXXDW
DT
     Patent
     English
LA
     ICM C07K005-08
IC
     ICS
          C07K005-10; C07K015-00; A61K037-02; A61K047-48; C08F289-00;
          C07K009-00; C08G069-10
     34-3 (Amino Acids, Peptides, and Proteins)
CC
     Section cross-reference(s): 1
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                                                                    DATE
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     EP 503301
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                                19920916
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EP 503301
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     JP 1991-297482
                          Α
                                 19911113
     JP 1992-22799
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                          A3
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CLASS
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                        C08F289-00; C07K009-00; C08G069-10
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                        C07K0005-08 [ICM, 5]; C07K0005-10 [ICS, 5]; C07K0005-00
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                        [I,C*]; A61K0047-48 [I,A]; A61P0035-00 [I,C*];
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                        C08F289/00; C08G069/10; C07K005/08B1
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                         530/329.000; 530/330.000; 530/331.000
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                         A61K047/48R; C07K005/08B1; C07K005/10B; C07K009/00D;
                         C07K014/78; C08F289/00; C08G069/10
OS
     MARPAT 118:125076
AΒ
     Fibronectin cell adhesion peptide fragments H-Z-D- or -L-Arg-X-Asp-Y-OH (X
     = L- or D-Leu, D-Ile, L- or D-Nle, L- or D-Phe, D-phenylglycine, D-Ala; Z,
     Y = independently bond, amino acid residue, or peptide residue, composed
     of Gly, Ser, Thr, L- or D-Asp, Ala, D-Glu, Pro), derivs., pharmaceutically
     acceptable salts, and pharmaceutical compns. comprising them were prepared
     as agents for inhibiting tumor metastasis.
ST
     antitumor fibronectin peptide fragment; neoplasm inhibitor fibronectin
     peptide conjugate
     Neoplasm inhibitors
ΤТ
        (fibronectin cell adhesion peptide fragments and conjugates)
TΤ
     Fibronectins
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (conjugates, peptide fragments and, preparation and antitumor activity of)
     3459-18-5
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (acetylation of)
     920-46-7
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (amidation of, with alanine)
     108-55-4
TT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (amidation of, with aminophenyl glucopyranosamine derivative)
ΙT
     145880-89-3
                   145880-90-6
                                  145880-91-7 145880-92-8
     145880-95-1
                   145880-96-2
                                  145880-97-3
                                                 145881-91-0
                                                               145881-92-1
     145881-93-2
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (amidation of, with chitin derivs.)
ΙT
     145880-93-9
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (amidation of, with chitin or chondroitin sulfate derivs.)
TΤ
     145880-98-4
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (amidation of, with chondroitin sulfate)
                                                    144377-70-8, PEO Acid 4000
ΙT
     77-92-9
               585-84-2
                           4023-65-8
                                       26106-63-8
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (amidation of, with fibronectin cell adhesion peptide fragments)
ΙT
     107-95-9, \beta-Alanine
     RL: RCT (Reactant); RACT (Reactant or reagent)
         (amidation of, with methacryloyl chloride)
ΙT
     145880-99-5
                    145881-01-2
     RL: RCT (Reactant); RACT (Reactant or reagent)
         (amidation of, with succinylated chondroitin sulfate)
ΙT
     145881-00-1
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RL: RCT (Reactant); RACT (Reactant or reagent)
        (amidation of, with trimellitoylated chondroitin sulfate)
IT
     6290-05-7
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (butoxycarbonylation of or double amidation of, with protected
        iminodiacetic acid)
                 7699-38-9
IT
     7284-16-4
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (conversion of, to fibronectin cell adhesion peptide fragment
        conjugates)
IT
     56808-39-0P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (conversion of, to fibronectin cell adhesion peptide fragment
        conjugates)
ΙT
     23680-31-1·
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (esterification and amidation reactions of)
     4530-20-5
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (esterification of, and peptide coupling reactions of, in preparation of
        fibronectin cell adhesion peptide fragments)
ΙT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (esterification of, with benzyl bromide)
     79-41-4, reactions
ΙT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (peptide coupling of, in preparation of fibronectin cell adhesion peptide
        fragments)
     51219-19-3
                  68262-71-5
                               145881-13-6
IΤ
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (peptide coupling of, with tetrapeptide ester)
IT
     13139-15-6
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (peptide coupling of, with tripeptide ester)
                            13574-13-5
                                          13734-34-4
                                                        16937-99-8
                                                                     18942-49-9
     6404-28-0
                 7536-58-5
ΙT
     33125-05-2
                 55674-63-0
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (peptide coupling reactions of, in preparation of antitumor peptides)
ΙT
     35150-07-3
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (peptide coupling reactions of, in preparation of fibronectin cell adhesion
        peptide fragments)
     145881-15-8P
TT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and acetylation of)
ΙT
     145881-84-1P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and acidic deblocking of)
IT
     145881-64-7P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and addition of, with sodium hydrogen sulfite)
IT
     145899-62-3P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
         (preparation and amidation of, with aconitic acid)
IT
     145881-69-2P
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RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and amidation of, with aconitic acid tris(aspartic acid)
        derivative)
ΙT
     145881-57-8P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and amidation of, with citric acid)
ΙT
     69171-62-6P
                   145881-61-4P
                                  145881-65-8P
                                                  145881-76-1P
     145881-83-0P
                    145881-88-5P
                                   145881-89-6P
                                                   145881-90-9P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and amidation of, with fibronectin cell adhesion peptide
        fragments)
ΙT
     65907-85-9P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and amidation of, with glutaric anhydride)
ΙT
     145881-63-6P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and amidation of, with iminodiacetic acid-tetracarboxylic acid
        derivative)
IT
     145881-82-9P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and amidation of, with succinic anhydride)
ΙT
     145880-86-0P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and amidation of, with succinylated polyallylamine)
     145881-70-5P
IT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and amidation of, with triethylenetetramine)
     1398-61-4DP, Chitin, acylated with dicarboxylic acid derivs., amides with
IT
     fibronectin cell adhesion peptide fragments
                                                   7512-17-6DP, oligomers,
     sulfated, ethers with fibronectin cell adhesion peptide fragments
     9007-28-7DP, acylated with dicarboxylic acid derivs., amides with
     fibronectin cell adhesion peptide fragments
                                                    30551-89-4DP, succinylated,
     amides with fibronectin cell adhesion peptide fragments
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                    145880-74-6P
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                                                      145880-85-9DP, ethers with
     ethers with sulfated oligo(acetylglucosamine)
     sulfated oligo(acetylglucosamine)
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                                   145880-88-2DP, amides with
     succinylated polyallylamine
                                145880-93-9DP, amides with succinylated
     carboxymethylated chitin
     carboxymethylated chitin and succinylated chondroitin sulfate
     145880-95-1DP, amides with sulfated carboxymethylated chitin
     145880-99-5DP, amides with succinylated chondroitin sulfate
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                    145881-07-8P
     145881-06-7P
                                   145881-11-4P
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     145991-54-4P
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
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study, unclassified); SPN (Synthetic preparation); BIOL (Biological
     study); PREP (Preparation)
        (preparation and antitumor activity of)
ΙT
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                    145881-66-9P
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     (Reactant or reagent)
        (preparation and catalytic deesterification of)
ΙT
     145881-16-9P
                    145881-17-0P
                                   145881-58-9P
                                                   145881-59-0P
                                                                  145881-77-2P
     145899-58-7P
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                                                   145991-51-1P
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     146225-99-2P
                    146245-93-4P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and catalytic hydrogenolysis of)
ΙΤ
     13089-27-5P
                   145881-81-8P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and catalytic reduction of)
ΙT
     63024-02-2P
                   145881-18-1P
                                  145881-42-1P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and deblocking of, with trifluoroacetic acid)
ΙT
     145881-53-4P
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     (Reactant or reagent)
        (preparation and desilylation of, with hydrogen fluoride)
ΙT
     56074-20-5P
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     (Reactant or reagent)
        (preparation and double amidation of, with iminodiacetate)
     115692-25-6P
ΙT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and peptide coupling of, with aspartic acid derivative)
     145881-67-0P
IT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and peptide coupling of, with fibronectin cell adhesion peptide
        fragments)
ΙT
     145881-86-3P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and peptide coupling of, with glycine derivative)
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ΙT
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     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and peptide coupling of, with leucine derivative)
     90013-41-5P
ΙT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and peptide coupling reactions of, in preparation of antitumor
        peptides)
IT
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        (preparation and polymerization of)
ΙT
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     (Reactant or reagent)
        (preparation and reductive deesterification of)
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     (Reactant or reagent)
        (preparation and saponification of)
IT
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     (Reactant or reagent)
        (preparation and silylation of)
ΙT
     63464-05-1P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation and substitution of, with fibronectin cell adhesion peptide
        fragment)
IT
     145880-09-7DP, amides with carboxymethylated chitin and itaconylated
     carboxymethylated chitin
                               145880-10-0DP, amides with succinylated
    polyallylamine, maleylated chondroitin sulfate, and
    145880-21-3P
                            145880-23-5P
                                           145880-24-6P
     glycol
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     145880-82-6DP, ethers with sulfated oligo(acetylglucosamine)
    145880-84-8DP, ethers with sulfated oligo(acetylglucosamine)
                                                145880-89-3DP,
     145880-87-1DP, amides with polyallylamine
     amides with maleylated carboxymethylated chitin
                                                      145880-90-6DP, amides
    with phthaloylated carboxymethylated chitin
                                                  145880-91-7DP, amides with
    carboxymethylated chitin
                               145880-92-8DP, amides with carboxymethylated
             145880-94-ODP, amides with sulfated carboxymethylated chitin
     145880-96-2DP, amides with sulfated succinylated carboxymethylated chitin
     145880-97-3DP, amides with sulfated succinylated carboxymethylated chitin
     145880-98-4DP, amides with chondroitin sulfate
                                                     145881-00-1DP, amides
    with trimellitoylated chondroitin sulfate
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     succinylated chondroitin sulfate
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     study); PREP (Preparation)
        (preparation of, as antitumor agent)
ΙT
     145881-75-0P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, catalytic hydrogenolysis, and succinylation of)
IT
     145881-87-4P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and amidation of with aminophenyl glucopyranoside
        derivative)
ΙT
     145991-20-4P
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation, deblocking, and amidation of, with methacryloyl chloride)
ΙT
     145881-12-5P
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                                                                145881-31-8P
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                                   145881-34-1P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
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(Reactant or reagent)
        (preparation, deblocking, and catalytic hydrogenolysis of)
ΙT
     145881-41-0P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and peptide coupling of, with
        (carboxyethyl) methacrylamide)
ΙT
     145881-24-9P
                    145881-25-0P
                                   145881-26-1P
                                                   145881-27-2P
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     145881-45-4P
                    145881-72-7P
                                   145899-59-8P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
    (Reactant or reagent)
        (preparation, deblocking, and peptide coupling of, with arginine derivative)
     145881-09-0P
                    145881-21-6P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and peptide coupling of, with arginine derivs.)
     69871-79-0P
                   80963-10-6P
                                 130457-93-1P
ΙT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and peptide coupling of, with aspartic acid
        derivative)
ΙT
     145881-46-5P
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     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and peptide coupling of, with glycine derivative)
IT
     145881-44-3P
                    145881-71-6P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and peptide coupling of, with leucine derivative)
     145881-48-7P
ΙT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
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        (preparation, deblocking, and peptide coupling of, with methacrylic acid)
     145881-94-3P
ΙT
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and peptide coupling of, with polyethylene glycol
        carboxylic acid derivative)
IΤ
     79113-14-7P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and peptide coupling of, with serine derivative)
IT
     145881-10-3P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, deblocking, and peptide coupling reactions of)
ΙT
     59178-90-4P
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation, polymerization, and peptide coupling of, with fibronectin cell
        adhesion peptide fragments)
IT
     145881-80-7P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, saponification, sulfation, and catalytic deblocking of)
IT
     145881-54-5P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation, sulfation, and catalytic hydrogenolysis of)
ΙT
     145881-85-2P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT .
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(Reactant or reagent)
        (preparation, sulfation, saponification, and catalytic deblocking of)
IT
     30551-89-4
    RL: RCT (Reactant); RACT (Reactant or reagent)
        (sequential succinylation and amidation of, with fibronectin cell
       adhesion peptide fragments)
IT
     9004-74-4
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (substitution of, with cyanuric chloride)
IT
     72708-10-2
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        (substitution of, with fibronectin cell adhesion peptide fragment)
IT
    145881-74-9
    RL: RCT (Reactant); RACT (Reactant or reagent)
        (substitution of, with glucopyranosyl bromide)
    108-77-0
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        (substitution of, with polyethylene glycol monomethyl ether)
     21085-72-3
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        (succinylation by, of pentaethylenehexaamine)
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        (succinylation of, with succinic anhydride)
     2791-79-9 ·
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     RL: RCT (Reactant); RACT (Reactant or reagent)
        (triple amidation of, with aconitic acid)
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L8
ΑN
    1993:14861 CAPLUS
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DN
    Entered STN: 10 Jan 1993
ΕD
    Electrically conductive-polymer-coated metal particles
ΤI
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    Hosokawa, Hiroshi; Kamada, Kensuke
PΑ
    Mitsubishi Rayon Co., Ltd., Japan
SO '
     Eur. Pat. Appl., 10 pp.
    CODEN: EPXXDW
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     ICS C08G061-12
     76-2 (Electric Phenomena)
     Section cross-reference(s): 38
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                        B32B0005-16 [ICM,5]; B32B0015-02 [ICS,5]; B32B0021-02
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                 NCL
                        428/403.000; 428/328.000; 428/407.000
AB
     Fine metal particles coated with elec. conductive polymers are
     protected from oxidation and have improved handling properties.
     conductive polymer may contain a polymeric electrolyte as a dopant.
ST
     conductive polymer coated metal powder
IΤ
     Electric conductors.
        (polymer-coated metal powders as, preparation of)
                  9003-01-4P, Polyacrylic acid 25087-26-7P, Polymethacrylic
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     9002-98-6P
            25191-25-7P
                          26336-38-9P, Polyvinylamine
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                     50851-57-5P, Polystyrenesulfonic acid
     Polyallylamine
     144892-84-2P
     RL: PREP (Preparation)
        (metal powders coated with conductive polymers doped with, preparation of)
ΙT
     25067-54-3P; Polyfuran
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     Polythiophene
     RL: PREP (Preparation)
        (metal powders coated with, preparation of)
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RL: PREP (Preparation)
         (powders, coated with elec. conductive polymers, preparation of)
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     ANSWER 19 OF 19 CAPLUS COPYRIGHT 2007 ACS on STN
     1989:436285 CAPLUS
ΑN
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ΤI
     A polymer-protected colloidal metal dispersion, a colloidal
     metal complex, and its preparation
IN
     Hirai, Hidefumi; Komiyama, Makoto; Otaki, Michitaka
PA
SO
     Eur. Pat. Appl., 52 pp.
     CODEN: EPXXDW
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     ICS A61K047-00; C12N011-14; G01N033-532
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     9-14 (Biochemical Methods)
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424/001.290; 424/001.110; 424/009.400; 424/009.420;
427/214.000; 427/216.000; 427/221.000; 428/407.000;
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                         516/077.000; 516/097.000; 523/205.000; 523/206.000;
                         523/375.000
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                  IPCR
                         B01J0013-00 [I,C*]; B01J0013-00 [I,A]; B01F0017-52
                         [I,C*]; B01F0017-52 [I,A]; B01J0031-06 [I,C*];
                         B01J0031-06 [I,A]; B01J0035-00 [I,C*]; B01J0035-12.
                         [I,A]; C08K0003-00 [I,C*]; C08K0003-02 [I,A];
                         C08K0003-08 [I,A]; C08L0057-00 [I,C*]; C08L0057-00
 JP 01020265
                  IPCI
                         C08L0101-00 [ICM, 4]; B01J0013-00 [ICS, 4]; B01J0031-06
                         [ICS, 4]; C08K0003-08 [ICS, 4]; C08K0003-00 [ICS, 4, C*];
                         A61K0043-00 [ICA,4]; A61K0049-02 [ICA,4]
                  IPCR
                         A61K0051-00 [I,C*]; A61K0051-00 [I,A]; B01J0013-00
                         [I,C*]; B01J0013-00 [I,A]; B01J0031-06 [I,C*];
                         B01J0031-06 [I,A]; C08F0008-00 [I,C*]; C08F0008-30
                         [I,A]; C08F0008-32 [I,A]; C08K0003-00 [I,C*];
                         C08K0003-02 [I,A]; C08K0003-08 [I,A]; C08L0101-00
                         [I,C*]; C08L0101-00 [I,A]
 US 5187209
                  IPCI
                         A61K0047-00 [ICM,5]; C08K0009-10 [ICS,5]; C08K0009-00
                         [ICS,5,C*]; C12N0011-14 [ICS,5]; C12N0011-00 [ICS,5,C*]
                  IPCR
                         A61K0009-51 [I,C*]; A61K0009-51 [I,A]; A61K0049-00
                         [I,C*]; A61K0049-00 [I,A]; B01J0013-00 [I,C*];
                         B01J0013-00 [I,A]; B01J0035-00 [N,C*]; B01J0035-00
                         [N,A]; C12N0015-02 [I,C*]; C12N0015-02 [I,A];
                         C12Q0001-68 [I,C*]; C12Q0001-68 [I,A]
                  NCL
                         424/001.290; 427/214.000; 427/216.000; 427/221.000;
                         428/403.000; 428/407.000; 516/077.000; 516/078.000;
                        -516/097.000; 523/205.000; 523/206.000; 523/375.000;
                         524/501.000
AΒ
     A polymer-protected colloidal metal dispersion comprises (a) a
     dispersion medium, (b) colloidal particles of Group Ib, VIIb, and VIII
     metals; and (c) a protective polymer including hydrazide, acrylic, and
     acrylamide polymers adsorbed on the colloidal particles. The colloidal
     metal dispersion is highly stable and the colloidal metal particles
     protected by the polymers can be easily and strongly bound to
     various amino group-containing compds. to give stable colloidal metal
     complexes. Acrylamide-N-vinyl-2-pyrrolidone copolymer containing 0.38 mequiv
     hydrazide groups/g (mol. weight .apprx.28,000) was dissolved in EtOH/H2O
     (1:1) and mixed with K2PtCl4. The mixture was irradiated with light from a
     500-W high-pressure Hg lamp to obtain a uniform blackish-brown dispersion
     of colloidal Pt protected by the hydrazide group-containing polymer.
     This polymer-protected colloidal Pt was conjugated to rabbit
     troponin via glutaraldehyde and the complex was reacted with paracryst.
     rabbit tropomyosin. The tropomyosin-labeled troponin complex was observed by
     electron microscopy.
ST
     colloidal metal dispersion polymer protection; platinum colloid hydrazide
     polymer protection; troponin tropomyocin colloidal platinum complex
IT
     Polyhydrazides
     Polymers, uses and miscellaneous
     RL: ANST (Analytical study)
         (colloidal metals protected by, proteins and other
        amine-containing compds. labeling with)
     Group IB elements
     Group VIIB elements
```

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Group VIII elements
     Metals, reactions
     RL: ANST (Analytical study)
        (colloidal, polymer-protected, proteins and other
        amine-containing compds. labeling with)
IT
     Hydrogenation catalysts
        (polymer-protected colloidal rhodium or platinum complexes
        with amino group-containing resins)
ΙT
     Vinyl compounds, compounds
     RL: ANST (Analytical study)
        (amino, complexes, with polymer-protected colloidal metals)
ΙT
     Nucleic acids
     Proteins, specific or class
     RL: ANST (Analytical study)
        (complexes, with polymer-protected colloidal metals)
ΙT
     Troponins
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (complexes, with polymer-protected colloidal platinum and
        tropomyosin, preparation of)
IT
     Albumins, compounds
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (complexes, with polymer-protected colloidal silver, preparation
        of)
IT
     Tropomyosins
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (complexes, with troponin labeled with polymer-protected
        colloidal platinum, preparation of)
TT
     79-10-7D, 2-Propenoic acid, esters, polymers
     RL: ANST (Analytical study)
        (colloidal metals protected by, proteins and other
        amine-containing compds. labeling with)
ΙT
     7439-88-5, Iridium, reactions
                                     7440-02-0, Nickel, reactions
                                                                    7440-04-2,
     Osmium, reactions
                        7440-05-3, Palladium, reactions 7440-06-4, Platinum,
     reactions
                 7440-15-5, Rhenium, reactions
                                                 7440-16-6, Rhodium, reactions
                                       7440-22-4, Silver, reactions
     7440-18-8, Ruthenium, reactions
                                    7440-57-5, Gold, reactions
     7440-50-8, Copper, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (colloidal, polymer-protected, proteins and other
        amine-containing compds. labeling with)
IT
     110-83-8, Cyclohexene, biological studies
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (hydrogenation of, by Diaion complexes with polymer-protected
        colloidal platinum catalyst)
IΤ
     107-13-1, Acrylonitrile, reactions
                                         109-92-2, Ethyl vinyl ether
     141-79-7, Mesityl oxide
                              592-41-6, 1-Hexene, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (hydrogenation of, polymer-protected colloidal
        rhodium-aminoethyl Bio-gel P-150 complex catalyst for)
     9003-05-8DP, Polyacrylamide, aminoalkylated, polymer-protected
ΤТ
                                9036-88-8DP, Mannan, aminated, polymer-
     colloidal metal complexes
     protected colloidal gold or platinum complexes 9060-90-6DP,
     Polyaminostyrene, polymer-protected colloidal metal complexes
     11028-71-0DP, Concanavalin A, polymer-protected colloidal gold
                30551-89-4P, Polyallylamine
     complexes
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of)
IT
     26124-23-2DP, Acrylamide-N-vinyl-2-pyrrolidone copolymer, hydrazine or
     azide reaction products
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of and colloidal gold or platinum protection by)
```

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IT
     119554-16-4DP, hydrazine or azide reaction products
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of and colloidal metal protection by)
     64387-83-3P, Acrylamide-methyl vinyl ketone copolymer
ΙT
                                                              76188-67-5P
                    119554-18-6DP, hydrazine or azide reaction products
     119554-14-2P
     119559-71-6DP, polymers, hydrazine reaction products
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of and colloidal platinum protection by)
IT
     25085-37-4P
                   25655-01-0P
                                 42936-66-3DP, Methacrylic acid-methyl vinyl
     ether copolymer, 4-nitrophenyl ester group-containing
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of and colloidal rhodium protection by)
     119554-13-1P
                    119554-15-3P
TΤ
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of and colloidal silver protection by)
     9047-06-7DP, Bio-Gel P 150, aminoethyl derivs., polymer-protected
TΤ
     colloidal rhodium complexes
                                   62862-86-6DP, Diaion, polymer-
     protected colloidal platinum complexes
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of, as hydrogenation catalysts)
IT
     16903-35-8, Chloroauric acid
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction of, in preparation of polymer-protected colloidal gold)
IT
     10025-99-7, Potassium tetrachloroplatinate
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction of, in preparation of polymer-protected colloidal
        platinum for troponin labeling)
IΤ
     12648-57-6, Rhodium chloride
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction of, in preparation of polymer-protected colloidal
        rhodium)
ΙT
     7761-88-8, Silver nitrate, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction of, in preparation of polymer-protected colloidal
        silver)
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- Dicarbonic acid, bis(1,1-dimethylethyl) ester (9CI)
- Formic acid, oxydi-, di-tert-butyl ester (7CI, 8CI)

OTHER NAMES:

- Bis(1,1-dimethylethyl) dicarbonate CN
- CN Bis(tert-butyl) dicarbonate
- CN BOC-anhydride
- CN Di-tert-butyl dicarbonate
- CN Di-tert-butyl oxydiformate
- CN Di-tert-butyl pyrocarbonate
- CN Pyrocarbonic acid di-tert-butyl ester
- CN tert-Butoxycarbonyl anhydride
- CN tert-Butyl dicarbonate
- MF C10 H18 O5
- CI COM
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9 POLYALLYLAMINE

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L11 0 AMINE AND L9

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=> s 19

L13 5391 L9

=> s 19 and allylamine 5391 L9 7884 ALLYLAMINE

L14 76 L9 AND ALLYLAMINE

=> s 114 and water 2550504 WATER

L15 . 6 L14 AND WATER

=> s 114 and(aq or water) 1086310 AQ 2550504 WATER

L16 9 L14 AND(AQ OR WATER)

=> d all 1-9

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TI Molecular Basis for Water-Promoted Supramolecular Chirality Inversion in Helical Rosette Nanotubes

AU Johnson, Ross S.; Yamazaki, Takeshi; Kovalenko, Andriy; Fenniri, Hicham

CS National Institute for Nanotechnology, Department of Chemistry and Department of Mechanical Engineering, University of Alberta, Edmonton, AB, T6G 2M9, Can.

SO Journal of the American Chemical Society (2007), 129(17), 5735-5743 CODEN: JACSAT; ISSN: 0002-7863

PB American Chemical Society

DT Journal

LA English

CC 22-3 (Physical Organic Chemistry) Section cross-reference(s): 3, 26, 34

AB Helical rosette nanotubes (RNTs) are obtained through the self-assembly of the G-C motif, a self-complementary DNA base analog featuring the complementary hydrogen bonding arrays of both guanine and cytosine. The first step of this process is the formation of a 6-membered supermacrocycle (rosette) maintained by 18 hydrogen bonds, which then self-organizes into a helical stack defining a supramol. sextuple helix whose chirality and three-dimensional organization arise from the chirality, chemical structure, and conformational organization of the G-C motif. Because a chiral G-C motif is predisposed to express itself asym. upon self-assembly, there is a natural tendency for it to form one chiral RNT over its mirror image. Here the authors describe the synthesis and characterization of a chiral G-C motif that self-assembles into helical

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RNTs in methanol, but undergoes mirror image supramol. chirality inversion
upon the addition of very small amts. of water (<1% volume/volume).
Extensive phys. and computational studies established that the
mirror-image RNTs obtained, referred to as chiromers, result from
thermodn. (in water) and kinetic (in methanol) self-assembly
processes involving two conformational isomers of the parent G-C motif.
Although derived from conformational states, the chiromers are
thermodynamically stable supramol. species, they display
dominant/recessive behavior, they memorize and amplify their chirality in
an achiral environment, they change their chirality in response to solvent
and temperature, and they catalytically transfer their chirality. A detailed
mechanism for supramol. chirality inversion triggered by specific mol.
interactions between water mols. and the G-C motif is proposed.
water promoted supramol chirality inversion helical rosette
nanotube
INDEXING IN PROGRESS
Catalysts
   (chirality inversion and transfer; mol. basis for water
   -promoted supramol. chirality inversion in helical rosette nanotubes)
Hydrogen bond
   (complimentary; mol. basis for water-promoted supramol.
   chirality inversion in helical rosette nanotubes)
Nanotubes
   (helical rosette; mol. basis for water-promoted supramol.
   chirality inversion in helical rosette nanotubes)
Chemical chains
   (helical; mol. basis for water-promoted supramol. chirality
   inversion in helical rosette nanotubes)
Chirality
   (inversion of supramol.; mol. basis for water-promoted
   supramol. chirality inversion in helical rosette nanotubes)
Atomic force microscopy
Helix (conformation)
NMR (nuclear magnetic resonance)
Overhauser spectroscopy
Rotamers
Scanning electron microscopy
Self-assembly
Self-association
Solvent effect
Supramolecular structure
Transmission electron microscopy
   (mol. basis for water-promoted supramol. chirality inversion
   in helical rosette nanotubes)
Nucleic acid bases
RL: PEP (Physical, engineering or chemical process); PRP (Properties);
PROC (Process); PRP (Properties)
   (mol. basis for water-promoted supramol. chirality inversion
  in helical rosette nanotubes)
Circular dichroism
   (normal, time-, and temperature dependent; mol. basis for water
   -promoted supramol. chirality inversion in helical rosette nanotubes)
67-56-1, Methanol 7732-18-5, Water
RL: CAT (Catalyst use); NUU (Other use, unclassified); USES (Uses)
   (mol. basis for water-promoted supramol. chirality inversion
   in helical rosette nanotubes)
68-12-2, DMF
RL: NUU (Other use; unclassified); RCT (Reactant); RACT (Reactant or
reagent); USES (Uses)
   (mol. basis for water-promoted supramol. chirality inversion
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in helical rosette nanotubes)
 ΙT
      67-52-7, Barbituric acid
                               100-51-6, Benzyl alcohol
      ALLYLamine
                  1336-21-6, Ammonium hydroxide
                                                   5470-11-1,
      Hydroxylamine hydrochloride
                                   10025-87-3, Phosphorus oxychloride
      20816-12-0, Osmium tetraoxide 24424-99-5
                                                 27738-96-1,
      Carbonisocyanatidic chloride
      RL: RCT (Reactant); RACT (Reactant or reagent)
         (mol. basis for water-promoted supramol. chirality inversion
         in helical rosette nanotubes)
 IT
      407-25-0, Trifluoroacetic anhydride
                                            7529-22-8, N-Methylmorpholine
     N-oxide
                7790-28-5, Sodium periodate
      RL: RGT (Reagent); RACT (Reactant or reagent)
         (mol. basis for water-promoted supramol. chirality inversion
         in helical rosette nanotubes)
 RE.CNT
               THERE ARE 83 CITED REFERENCES AVAILABLE FOR THIS RECORD
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- DN 146:229612
- Entered STN: 09 Feb 2007 ·ED
- Preparation of macrocyclic carboxylic acids, amides, and acylsulfonamides ΤI as inhibitors of HCV replication
- ΙN Seiwert, Scott D.; Blatt, Lawrence M.; Andrews, Steven W.; Martin, Pierre; Schumacher, Andreas; Barnett, Bradley R.; Eary, Todd C.; Kaus, Robert; Kercher, Timothy; Liu, Weidong; Lyon, Michael; Nichols, Paul; Wang, Bin; Sammakia, Tarek; Kennedy, April; Jiang, Yutong
- Intermune, Inc., USA; Array Biopharma Inc. PA
- PCT Int. Appl., 512pp. SO CODEN: PIXXD2
- DT Patent
- English
- 34-3 (Amino Acids, Peptides, and Proteins) Section cross-reference(s): 1, 7, 63

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                             [I,A]; C07C0231-00 [I,C*]; C07C0269-00 [I,A];
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      CASREACT 146:229612; MARPAT 146:229612
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- * STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY AVAILABLE VIA OFFLINE PRINT *
- The invention relates to macrocyclic compds. I and analogs [R1 = H, OC(:O)R1; R1 = (un)substituted N-heteroaryl; R2 = OH, NHR5; R5 = Ph, alkyl, CN, cyclopropylcarbonyl, etc.; R3 = H, CH2R6, CSNH2, (un)substituted thiazol-2-yl, etc.; R6 = CF3, t-Bu, (un)substituted Ph, cyclopropyl, furanyl, etc.; R4 = H, cyclopropylmethyl; the dashed line represents an optional double bond], and their pharmaceutically acceptable salts, prodrugs, and esters for use in pharmaceutical compns. for the treatment of hepatitis C virus (HCV) infection and liver fibrosis. Thus, compound II, prepared by reaction of the macrocyclic prolinol derivative with

cDI in the presence of DCE and treatment with 1-methylcyclopropane-1-sulfonamide in the presence of DBU, showed IC50 < 0.1 μ M in the NS3-NS4 protease inhibition assay.

ST peptide macrocyclic carboxylic acid amide acylsulfonamide prepn inhibitor

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HCV
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     Peptides, preparation
     RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
     (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
        (cyclic; preparation of macrocyclic carboxylic acids, amides and
        acylsulfonamides as inhibitors of HCV replication)
ΙT
    Liver, disease
        (fibrosis; preparation of macrocyclic carboxylic acids, amides and
        acylsulfonamides as inhibitors of HCV replication)
     Fibrosis
IT
        (hepatic; preparation of macrocyclic carboxylic acids, amides and
        acylsulfonamides as inhibitors of HCV replication)
ΙT
     Drug delivery systems
        (injections, s.c.; preparation of macrocyclic carboxylic acids, amides and
        acylsulfonamides as inhibitors of HCV replication)
ΙΤ
     Antiviral agents
     Hepatitis C virus
     Human
        (preparation of macrocyclic carboxylic acids, amides and acylsulfonamides as
        inhibitors of HCV replication)
IT
    Macrocyclic compounds
     RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic
     preparation); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); RACT (Reactant or reagent); USES (Uses)
        (preparation of macrocyclic carboxylic acids, amides and acylsulfonamides as
        inhibitors of HCV replication)
IT
     Polyoxyalkylenes, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (preparation of macrocyclic carboxylic acids, amides and acylsulfonamides as
        inhibitors of HCV replication)
IT
     Infection
        (viral; preparation of macrocyclic carboxylic acids, amides and .
        acylsulfonamides as inhibitors of HCV replication)
IT
     Interferons
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (a; preparation of macrocyclic carboxylic acids, amides and
        acylsulfonamides as inhibitors of HCV replication)
ΙT
     Interferons
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (y; preparation of macrocyclic carboxylic acids, amides and
        acylsulfonamides as inhibitors of HCV replication)
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     394730-60-0, SCH 503034
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        (NS3 protease inhibitor; combination therapy agent; preparation of
        macrocyclic carboxylic acids, amides, and acylsulfonamides as
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                 7481-89-2, 2' 3' Dideoxycytidine
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                      36791-04-5, Ribavirin
                                              53179-13-8D, Pirfenidone, analogs
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     69655-05-6, 2' 3' Dideoxyinosine 113852-37-2, Cidofovir 119567-79-2,
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                                            136470-78-5, Abacavir
     Viramidine
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     142340-99-6
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     170277-31-3, Infliximab
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     331731-18-1, Adalimumab 518974-95-3, Thymosin \alpha
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        (combination therapy agent; preparation of macrocyclic carboxylic acids,
        amides and acylsulfonamides as inhibitors of HCV replication)
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Ι·Τ
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
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(conjugates with IFN- α ; combination therapy agents; preparation of

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macrocyclic carboxylic acids, amides and acylsulfonamides as inhibitors
        of HCV replication)
IT
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     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (conjugates with monoPEG; combination therapy agents; preparation of
        macrocyclic carboxylic acids, amides and acylsulfonamides as inhibitors
        of HCV replication)
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                 9028-93-7
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        (inhibitors; combination therapy agents; preparation of macrocyclic
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        replication)
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     149885-80-3, Ns3 ns4a protease
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     (Reactant or reagent)
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        (preparation of macrocyclic carboxylic acids, amides and acylsulfonamides as
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     5-(Hydroxymethyl) furan-2-carboxaldehyde
                                              67-56-1, Methanol, reactions
     70-11-1, 2-Bromo-1-phenylethanone 75-03-6, Iodoethane
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     2-Methylpropan-1-amine 90-04-0, 2-Methoxyaniline
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     2-Chloroaniline 95-76-1, 3,4-Dichloroaniline
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     98-64-6, 4-Chlorobenzenesulfonamide 100-61-8, N-Methyl-N-phenylamine,
                 103-67-3, N-Benzylmethylamine 103-82-2D, 2-Phenylacetic
     acid, resin-bound 104-01-8D, 2-(4-Methoxyphenyl)acetic acid, resin-bound
     104-86-9
                106-49-0, p-Toluidine, reactions
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                107-11-9, Allylamine
     N, N-Methyl-1, 2-ethanediamine
                                   108-37-2, m-Chlorophenyl bromide
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     117-21-5, 3-Chlorophthalic anhydride
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                                   319-03-9, 4-Fluorophthalic anhydride
     136-95-8, 2-Benzothiazolamine
     371-40-4, 4-Fluoroaniline
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     402-46-0, 4-Fluorobenzenesulfonamide
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     (Trifluoromethyl) benzaldehyde 459-59-6 460-40-2, 3,3,3-
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             701-34-8, 4-Bromobenzenesulfonamide 714-15-8
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     1,4-Dichloro-2-butene 765-30-0, Cyclopropylamine
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     (E)-1,4-Dibromobut-2-ene 872-53-7, Cyclopentanecarboxaldehyde
     1070-83-3D, 3,3-Dimethylbutanoic acid, resin-bound 1123-56-4,
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     2,6-Dimethylbenzaldehyde
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1489-69-6, Cyclopropylcarboxaldehyde 1524-40-9, 3-
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Fluorobenzenesulfonamide
1694-92-4, 2-Nitrobenzene-1-sulfonyl chloride
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6,7-Dimethoxy-1,2,3,4-tetrahydroisoquinoline
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2-(4-Chlorophenyl)acetic acid, resin-bound
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4-Morpholineethanamine
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3-(Aminomethyl)benzoic acid
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Cyclohexanesulfonamide
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[(Cyclopropyl)methyl]amine
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2835-77-0, 2-Aminobenzophenone 3118-68-1, 3-Cyanobenzenesulfonamide 3119-02-6, 4-Cyanobenzenesulfonamide 3144-04-5, 1-Butanesulfonamide 3731-51-9, 2-(Aminomethyl)pyridine 3731-53-1, 4-(Aminomethyl)pyridine
3886-69-9, (R)-1-Phenylethanamine 3984-17-6 3984-18-7, N-Cyclohexylsulfamide 4108-90-1, 1-Piperidinesulfonamide
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4-Bromobut-1-ene
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Cyclopropanecarboxamide 6325-93-5, 4-Nitrobenzenesulfonamide
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7103-09-5, But-3-enylmagnesium bromide 7647-01-0, Hydrochloric acid,
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                                              7681-84-7,
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Tetrahydrofuran-2-carboxaldehyde
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                                          13726-69-7, (2S,4R)-1-(tert-
(4-Methoxyphenyl)sulfamide
Butoxycarbonyl)-4-hydroxypyrrolidine-2-carboxylic acid
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15761-39-4, (S)-1-(tert-Butoxycarbonyl)pyrrolidine-2-carboxylic acid
15959-53-2, Phenylsulfamide 16993-47-8, 2-Phenylethanesulfonamide
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Dichlorobenzenesulfonamide 20532-15-4, 2,4-Dichlorobenzenesulfonamide
21508-19-0, 5-Chlorofuran-2-carboxaldehyde 22884-29-3,
Isobutyltriphenylphosphonium bromide 24243-71-8, Propanesulfonamide
24424-99-5, Boc anhydride 26238-14-2, 4-Trifluoromethylphthalic
anhydride 27578-60-5, 1-Piperidineethanamine 27757-85-3,
(Thiophen-2-y1) methanamine 27854-96-2, (S)-1-(Pyridin-4-y1) ethylamine
28460-30-2 30058-40-3, 2-Fluorobenzenesulfonamide 31795-93-4,
N-Benzyl-L-proline 32085-88-4, 3,5-Difluorobenzaldehyde 32327-43-8,
5-Chloronaphthalene-1-sulfonamide 32857-62-8D, resin-bound
34036-07-2, 3,4-Difluorobenzaldehyde 34813-49-5, tert-Butylsulfonamide
35203-92-0 45159-34-0
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50715-28-1, Cyclopentyl chloroformate . 51108-29-3
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4,5-Dimethylfuran-2-carboxaldehyde
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Methoxybenzenesulfonamide
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3-Methoxybenzenesulfonamide
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tetrahydroisoguinoline
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tetrahydroisoquinoline 89556-74-1
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Bromobenzenesulfonamide
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98198-68-6
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139631-62-2, Cyclopropanesulfonyl chloride 147962-41-2
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N-Ethyl-N-methylsulfamide
                            164648-70-8, (4-Cyanophenyl) sulfamide
               175137-46-9, 3-Amino-5-cyclopropyl-1H-pyrazole 176673-04-4
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(R)-2-Methylpropane-2-sulfinamide 205487-11-2
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     Octahydropyrimido[1,2-a]azepine
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     16940-66-2, Sodium borohydride
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        as inhibitors of HCV replication)
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     2006:1155411 CAPLUS
ΑN
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ED
     Entered STN: 03 Nov 2006
TΙ
     Preparation of piperidine derivatives as tachykinin receptor antagonists
IN
     Nagaoka, Naomi; Marunaka, Shigeyuki; Fukuta, Makoto
     Takeda Pharmaceutical Company Limited, Japan
PA
SO
     PCT Int. Appl., 323pp.
     CODEN: PIXXD2
DT
     Patent
     Japanese
LΑ
     28-10 (Heterocyclic Compounds (More Than One Hetero Atom))
CC
     Section cross-reference(s): 1, 27, 63
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          RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
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PRAI JP 2005-124335
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CLASS
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                           A61K009/20H6F2; A61K009/20H4B
                   ECLA
OS
     MARPAT 145:471540
AB
     The title compds. (no biol. data) are prepared This document discloses a
     pharmaceutical composition comprising N-(2-[(3R,4S)-4-((2-methoxy-5-[5-
      (trifluoromethyl)-1H-tetrazol-1-yl]benzyl)amino)-3-phenylpiperidin-1-yl]-2-
      oxoethyl)acetamide (I), a salt or a prodrug thereof, a sugar and a
     hydrophilic water-insol. substance. Thus, N-(2-[(3R,4S)-4-((2-
     hydroxy-5-[5-(trifluoromethyl)-1H-tetrazol-1-yl]benzyl)amino)-3-
      phenylpiperidin-1-yl]-2-oxoethyl)acetamide was prepared in 3 steps from
      (3R,4S)-4-amino-3-phenylpiperidine-1-carboxylic acid tert-Bu ester and
      2-hydroxy-5-[5-(trifluoromethyl)-1H-tetrazol-1-yl]benzaldehyde.
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Formulations containing I are given. Tablets containing I showed high elution
     stability.
     piperidine deriv tachykinin receptor antagonist prepn; pharmaceutical
ST
     compn piperidine deriv tachykinin receptor antagonist prepn
ΙT
     Bladder, disease
     Inflammation
        (cystitis, interstitial; preparation and use of piperidine derivs. as
        tachykinin receptor antagonists) .
ΙT
     Mental and behavioral disorders
        (depression; preparation and use of piperidine derivs. as tachykinin
        receptor antagonists)
ΙΤ
     Disease, animal
        (digestion disorder; preparation and use of piperidine derivs. as tachykinin
        receptor antagonists)
ΙT
     Digestion, biological
        (disease; preparation and use of piperidine derivs. as tachykinin receptor
        antagonists)
IT
     Stability
        (elution stability; of pharmaceutical composition comprising piperidine
        derivative, salt or prodrug thereof, sugar and hydrophilic water
        -insol. substance)
IT
     Granulation
        (fluid layer; pharmaceutical composition comprising piperidine derivative,
salt
        or prodrug thereof, sugar and hydrophilic water-insol.
        substance)
IT.
     Intestine, disease
        (inflammatory; preparation and use of piperidine derivs. as tachykinin
        receptor antagonists)
TT
     Intestine, disease
        (irritable bowel syndrome; preparation and use of piperidine derivs. as
        tachykinin receptor antagonists)
ΙT
     Urinary system, disease
        (lower urinary tract; preparation and use of piperidine derivs. as
        tachykinin receptor antagonists)
İΤ
     Mental and behavioral disorders
        (neurosis; preparation and use of piperidine derivs. as tachykinin receptor
        antagonists)
ΙT
     Crystal structure
        (of piperidine derivative)
IT
        (overactive; preparation and use of piperidine derivs. as tachykinin
        receptor antagonists)
ΙT
        (pelvic; preparation and use of piperidine derivs, as tachykinin receptor
        antagonists)
     Granulation
ΙT
        (pharmaceutical composition comprising piperidine derivative, salt or
prodrug
        thereof, sugar and hydrophilic water-insol. substance)
ΙT
     Alditols
     Carbohydrates, biological studies
     Disaccharides
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses).
        (pharmaceutical composition comprising piperidine derivative, salt or
prodrug
        thereof, sugar and hydrophilic water-insol. substance)
     Salts, preparation
     RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
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(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (pharmaceutical composition comprising piperidine derivative, salt or
prodrug
        thereof, sugar and hydrophilic water-insol. substance)
ΙT
     Analgesics
     Anti-inflammatory agents
     Antidepressants
     Antiemetics
     Anxiety
     Anxiolytics
     Central nervous system, disease
     Central nervous system agents
     Hypnotics and Sedatives
     Insomnia
     Vomiting
        (preparation and use of piperidine derivs. as tachykinin receptor
        antagonists)
ΙT
     Tachykinin receptors
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (preparation of piperidine derivs. as tachykinin receptor antagonists)
ΙT
     Drug delivery systems
        (prodrugs; pharmaceutical composition comprising piperidine derivative,
salt or
        prodrug thereof, sugar and hydrophilic water-insol.
        substance)
ΙT
     Drug delivery systems
        (solids; pharmaceutical composition comprising piperidine derivative, salt
or
        prodrug thereof, sugar and hydrophilic water-insol.
        substance)
TT
     Drug delivery systems
        (tablets; pharmaceutical composition comprising piperidine derivative, salt
or
        prodrug thereof, sugar and hydrophilic water-insol.
        substance)
ΙT
     Granulation
        (wet; pharmaceutical composition comprising piperidine derivative, salt or
        prodrug thereof, sugar and hydrophilic water-insol.
        substance)
     9004-64-2, Hydroxypropylcellulose
TΤ
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (low substituted; pharmaceutical composition comprising piperidine
derivative,
        salt or prodrug thereof, sugar and hydrophilic water-insol.
        substance)
IT
     9004-34-6, Cellulose, biological studies
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BİOL
     (Biological study); USES (Uses)
        (microcryst.; pharmaceutical composition comprising piperidine derivative,
salt
        or prodrug thereof, sugar and hydrophilic water-insol.
        substance)
IΤ
     69-65-8, Mannitol
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (pharmaceutical composition comprising piperidine derivative, salt or
        thereof, sugar and hydrophilic water-insol. substance)
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632344-29-7P, cis-4-[[3,5-Bis(trifluoromethyl)benzyl]oxy]-N-methyl-3-
       phenyl-1-piperidinecarboxamide
                                                            632345-06-3P
                                                                                    632350-58-4P
       632352-48-8P
                                632352-49-9P
                                                       632352-50-2P
                                                                               632352-51-3P
       RL: ANT (Analyte); PAC (Pharmacological activity); SPN (Synthetic
       preparation); THU (Therapeutic use); ANST (Analytical study); BIOL
        (Biological study); PREP (Preparation); USES (Uses)
             (preparation of piperidine derivs. as tachykinin receptor antagonists)
IT
        913092-51-0P
                                913092-52-1P
       RL: ANT (Analyte); RCT (Reactant); SPN (Synthetic preparation); ANST
        (Analytical study); PREP (Preparation); RACT (Reactant or reagent)
             (preparation of piperidine derivs. as tachykinin receptor antagonists)
IT
        632352-59-1P, trans-4-[(2-Methoxybenzyl)amino]-3-phenylpiperidine-1-
        carboxylic acid tert-butyl ester
       RL: BYP (Byproduct); PREP (Preparation)
             (preparation of piperidine derivs. as tachykinin receptor antagonists)
        913092-71-4P
TT
        RL: PAC (Pharmacological activity); PRP (Properties); PUR (Purification or
        recovery); SPN (Synthetic preparation); THU (Therapeutic use); BIOL
        (Biological study); PREP (Preparation); USES (Uses)
             (preparation of piperidine derivs. as tachykinin receptor antagonists)
ΙT
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        790657-29-3P
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        913168-48-6P
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        913976-53-1P
                                913976-60-0P
                                                        913976-61-1P
        RL: PAC (Pharmacological activity); PUR (Purification or recovery); SPN
        (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study);
        PREP (Preparation); USES (Uses)
             (preparation of piperidine derivs. as tachykinin receptor antagonists)
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        cis-4-[[3-Nitro-5-(trifluoromethyl)benzyl]oxy]-3-phenylpiperidine
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        RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic
        preparation); THU (Therapeutic use); BIOL (Biological study); PREP
        (Preparation); RACT (Reactant or reagent); USES (Uses)
             (preparation of piperidine derivs. as tachykinin receptor antagonists)
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        (+) - (3R, 4S) - 1 - [(1-Acetyl-4-piperidinyl)carbonyl] - 4 - [[3, 5-piperidinyl]carbonyl] arbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl]carbonyl[carbonyl]carbonyl[carbonyl]carbonyl[carbonyl]carbonyl[carbonyl]carbonyl[carbonyl]carbonyl[carbonyl]carbonyl[carbonyl]carbonyl[carbonyl
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     RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
     (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
         (preparation of piperidine derivs. as tachykinin receptor antagonists)
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bis(trifluoromethyl)benzyl]oxy]-3-phenylpiperidine

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                                     632349-43-0P
                                                     632349-44-1P
     RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
     (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (preparation of piperidine derivs. as tachykinin receptor antagonists)
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                                              632351-77-0P
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632351-85-0P
                632351-86-1P
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632352-38-6P
                632352-39-7P
                               632352-40-0P
                                              632352-41-1P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
 (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
    (preparation of piperidine derivs. as tachykinin receptor antagonists)
632352-42-2P
                632352-43-3P
                               632352-44-4P
                                              632352-45-5P
                                                             632352-46-6P
 632352-47-7P
                913092-57-6P
                               913092-58-7P
                                              913092-59-8P
                                                             913092-60-1P
913092-61-2P
                913092-.62-3P
                               913092-63-4P
                                              913092-65-6P
                                                             913092-68-9P
913092-70-3P
                913976-54-2P
                               913976-55-3P
                                              913976-56-4P
                                                             913976-57-5P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
 (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
    (preparation of piperidine derivs. as tachykinin receptor antagonists)
.64-04-0, Phenethylamine 64-19-7, Acetic acid, reactions
                          75-36-5, Acetyl chloride
Methyl iodide, reactions
                                                       79-04-9,
                       79-09-4, Propionic acid, reactions
Chloroacetyl chloride
                                                              79-22-1,
Chloroformic acid methyl ester 79-44-7, N,N-Dimethylcarbamic acid
                                94-53-1, Piperonylic acid
           91-01-0, Benzhydrol
chloride
                                                             98-88-4,
Benzoyl chloride
                   100-39-0, Benzyl bromide 100-51-6, Benzyl alcohol,
reactions
            100-97-0, Hexamethylenetetramine, reactions
                                                           101-97-3,
Phenylacetic acid ethyl ester 103-71-9, Phenylisocyanate, reactions
105-36-2, Bromoacetic acid ethyl ester
                                         107-10-8, Propylamine, reactions
107-11-9, Allylamine 108-86-1, Bromobenzene, reactions
109-09-1, 2-Chloropyridine 109-90-0, Ethylisocyanate
                                                        123-38-6,
Propionaldehyde, reactions 124-63-0, Methylsulfonyl chloride 135-02-4,
                 402-49-3, 4-(Trifluoromethyl)benzyl bromide
o-Anisaldehyde
                                                               407-25-0,
Trifluoroacetic acid anhydride
                                 452-63-1, 4-Fluoro-2-methyl-1-
bromobenzene
                460-00-4, 4-Fluorobromobenzene
                                                 530-62-1,
N, N'-Carbonyldiimidazole
                           543-24-8, N-Acetylglycine
                                                        556-61-6,
                       590-17-0, Bromoacetonitrile
Methylisothiocyanate
                                                      612-16-8,
 2-Methoxybenzyl alcohol
                           624-83-9, Methylisocyanate
                                                        625-45-6,
Methoxyacetic acid
                      683-57-8, Bromoacetamide · 725-89-3,
 3,5-Bis(trifluoromethyl)benzoic acid
                                        753-90-2, 2,2,2-Trifluoroethylamine
 1939-99-7, \alpha-Toluenesulfonylchloride
                                        2450-71-7, Propargylamine
 2491-06-7, N,N-Dimethylglycine hydrochloride 2491-15-8, N-Formylglycine
2627-86-3, (S)-1-Phenylethylamine
                                   2949-22-6, Isocyanatoacetic acid ethyl
        2999-46-4, Ethylisocyanoacetate 3251-69-2, 4-Imidazoleacetic
acid hydrochloride 3886-69-9, (R)-1-Phenylethylamine
                                                         3886-70-2,
 (R) -1-Naphthylethylamine
                           4187-38-6, (R) -4-Tolylethylamine
 4530-20-5, Boc-glycine 4637-24-5, Dimethylformamide dimethylacetal
            6638-79-5, N,O-Dimethylhydroxylamine hydrochloride
 7693-46-1, Chloroformic acid 4-nitrophenyl ester
                                                   13750-81-7,
 1-Methyl-1H-imidazole-2-carboxaldehyde 13889-98-0, 1-Acetylpiperazine
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13939-69-0, 1-Piperidinecarbonyl chloride 15285-15-1,
     5-Chloro-3-methyl-1,2,4-triazole 15674-67-6, 3-Diethylaminopropionic
    acid hydrochloride 16179-97-8 24424-99-5, Di-tert-butyl
                   25016-01-7, 5-Bromo-2-methoxybenzaldehyde
    dicarbonate
                                                               25503-90-6,
     1-Acetylpiperidine-4-carboxylic acid 26371-07-3, 1-Piperidinepropionic
            32161-06-1, 1-Acetyl-4-piperidinone
                                                  32247-96-4,
     3,5-Bis(trifluoromethyl)benzyl bromide
                                             33524-31-1, 2,5-Dimethoxybenzyl
               42933-43-7, 2,3-Dihydro-1-benzofuran-5-amine
    alcohol
                                                              58936-19-9
     63327-49-1, (Acetylamino) (hydroxy) acetic acid
                                                     76041-06-0,
     3-[(3-Ethoxy-3-oxopropyl)amino]-2-phenylpropanoic acid ethyl ester
     76393-16-3, Ethyl 2-isocyanatobenzoate
                                             79099-07-3, 4-0xo-1-
    piperidinecarboxylic acid tert-butyl ester
                                                 84358-13-4
    159689-88-0, 3-(Trifluoromethoxy)benzyl bromide
                                                      168267-01-4,
     2-Hydroxy-5-[5-(trifluoromethyl)-1H-tetrazol-1-yl]benzaldehyde
    168267-11-6, 2-Methoxy-5-(5-(trifluoromethyl))-1H-tetrazol-1-
    yl)benzaldehyde
                       225246-36-6, 2-(Cyclopropyloxy)-5-[5-(trifluoromethyl)-
     1H-tetrazol-1-yl]benzaldehyde
                                    239087-09-3, 3-Fluoro-5-
     (trifluoromethyl)benzyl bromide 252742-72-6, 5-(Chloromethyl)-2,4-
    dihydro-3H-1,2,4-triazol-3-one
                                      304457-86-1, 1-(1H-Imidazol-4-yl)-1-
    propanone
                 632353-50-5, Methanesulfonic acid 3-nitro-5-
     (trifluoromethyl)benzyl ester 632353-52-7, cis-4-Amino-3-[bis(4-
     fluorophenyl)methyl]-1-piperidinecarboxylic acid tert-butyl ester
    880135-38-6, (3R,4S)-4-Amino-3-phenylpiperidine-1-carboxylic
    acidtert-butyl ester
                           913092-28-1
                                          913252-06-9, (3S,4R)-4-Amino-3-
    phenylpiperidine-1-carboxylic acid tert-butyl ester
    RL: RCT (Reactant); RACT (Reactant or reagent)
        (preparation of piperidine derivs. as tachykinin receptor antagonists)
     5779-95-3P, 3,5-Dimethylbenzaldehyde 76041-09-3P, 3-Phenyl-4-piperidone
IT
     183808-94-8P; 2-Ethoxy-5-[5-(trifluoromethyl)-1H-tetrazol-1-
    vl]benzaldehyde
                      193274-82-7P, 3-Benzyl-4-oxopiperidine-1-carboxylic acid
     tert-butyl ester
                      219324-21-7P
                                       444288-42-0P
                                                     446302-83-6P,
     1-Benzyl-3-phenyl-4-piperidinone
                                       562837-80-3P, 1-Acetyl-3-benzhydryl-4-
    piperidinone · 562837-99-4P
                                  632352-53<del>-</del>5P
                                                 632352-54-6P,
     3-[Benzyl(3-ethoxy-3-oxopropyl)amino]-2-phenylpropanoic acid ethyl ester
     632352-55-7P, 1-Benzyl-4-hydroxy-5-phenyl-1,2,5,6-tetrahydro-3-
    pyridinecarboxylic acid ethyl ester 632352-56-8P, 4-0xo-3-
    phenylpiperidine-1-carboxylic acid tert-butyl ester
                                                          632352-57-9P,
     4-Hydroxy-3-phenylpiperidine-1-carboxylic acid tert-butyl ester
     632352-58-0P, cis-4-[(2-Methoxybenzyl)amino]-3-phenylpiperidine-1-
    carboxylic acid tert-butyl ester 632352-60-4P, 4-Amino-3-
    phenylpiperidine-1-carboxylic acid tert-butyl ester 632352-61-5P,
    cis-3-Benzhydryl-4-[[3,5-bis(trifluoromethyl)benzyl]oxy]piperidine-1-
    carboxylic acid tert-butyl ester 632352-62-6P, 3-Benzhydryl-4-
     oxopiperidine-1-carboxylic acid tert-butyl ester
                                                       632352-63-7P,
     3-Benzhydryl-4-hydroxypiperidine-1-carboxylic acid tert-butyl ester
     632352-64-8P, cis-3-Benzhydryl-4-[[3-fluoro-5-
     (trifluoromethyl)benzyl]oxy]piperidine-1-carboxylic acid tert-butyl ester
     632352-65-9P, cis-3-Benzhydryl-4-[[3-(trifluoromethoxy)benzyl]oxy]piperidi
     ne-1-carboxylic acid tert-butyl ester 632352-66-0P, cis-3-Benzhydryl-4-
     [[4-(trifluoromethyl)benzyl]oxy]piperidine-1-carboxylic acid tert-butyl
             632352-67-1P, 5-[5-(Trifluoromethyl)-1H-tetrazol-1-yl]-2,3-dihydro-
     1-benzofuran-7-carboxaldehyde
                                    632352-68-2P, N-(2,3-Dihydro-benzofuran-5-
     yl)-2,2,2-trifluoro-acetamide
                                     632352-69-3P
                                                   632352-70-6P
                                                                  632352-71-7P
     632352-72-8P
                    632352-73-9P
                                   632352-74-0P, 3-(4-Fluorophenyl)-4-oxo-1-
    piperidinecarboxylic acid tert-butyl ester
                                                 632352-75-1P,
    cis-3-(4-Fluorophenyl)-4-hydroxy-1-piperidinecarboxylic acid tert-butyl
             632352-76-2P
                           632352-77-3P
                                           632352-78-4P
                                                         632352-79-5P
     632352-80-8P
                    632352-81-9P
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    632353-00-5P
                                   632353-02-7P, cis-4-[[2-Methoxy-5-[5-
                    632353-01-6P
     (trifluoromethyl)-1H-tetrazol-1-yl]benzyl]amino]-3-phenyl-1-
    piperidinecarboxylic acid tert-butyl ester
                                                  632353-03-8P
                                                                 632353-04-9P
                   632353-06-1P
                                   632353-07-2P
    632353-05-0P
                                                  632353-08-3P
                                                                 632353-09-4P
     632353-10-7P, cis-3-Phenyl-4-[[[5-[5-(trifluoromethyl)-1H-tetrazol-1-yl]-
    2,3-dihydro-1-benzofuran-7-yl]methyl]amino]-1-piperidinecarboxylic acid
                      632353-11-8P 632353-13-0P
    tert-butyl ester
                                                     632353-15-2P
     632353-17-4P, cis-3-Benzyl-4-[[3,5-bis(trifluoromethyl)benzyl]oxy]piperidi
    ne-1-carboxylic acid tert-butyl ester
                                             632353-21-0P, .cis-3-Benzyl-4-
    hydroxypiperidine-1-carboxylic acid tert-butyl ester
                                                            632353-23-2P
     632353-26-5P
                    632353-28-7P, cis-tert-Butyl 4-amino-3-phenylpiperidine-1-
                   632353-30-1P, cis-4-[[4-[[3,5-Bis(trifluoromethyl)benzyl]oxy
    carboxylate
     ]-3-phenyl-1-piperidinyl]carbonyl]-1-piperidinecarboxylic acid tert-butyl
             632353-32-3P, cis-4-[[3,5-Bis(trifluoromethyl)benzyl]oxy]-3-phenyl-
    1-(4-piperidinylcarbonyl)piperidine hydrochloride
                                                        632353-34-5P,
     (3R,4S)-4-Hydroxy-3-phenyl-1-piperidinecarboxylic acid tert-butyl ester
     632353-36-7P, (3R,4S)-4-[[3,5-Bis(trifluoromethyl)benzyl]oxy]-3-
    phenylpiperidine hydrochloride 632353-39-0P
                                                     632353-41-4P
     632353-42-5P, cis-4-[[(Benzyloxy)carbonyl]amino]-3-[bis(4-
     fluorophenyl)methyl]-1-piperidinecarboxylic acid tert-butyl ester
     632353-43-6P
                   632353-44-7P
                                   632353-45-8P, cis-4-Amino-3-[bis(4-
     fluorophenyl)methyl]-N-ethyl-1-piperidinecarboxamide hydrochloride
     632353-46-9P
                    632353-47-0P
                                   632353-48-1P, N-Ethyl-4-oxo-3-
                                      910875-38-6P
                                                     910875-39-7P,
     phenylpiperidine-1-carboxamide
     3-Phenylpiperidin-4-one hydrochloride
                                             913092-27-0P
                                                            913092-29-2P
     913092-30-5P
                                   913092-32-7P
                                                  913092-33-8P
                                                                  913092-34-9P
                    913092-31-6P
                                                  913092-38-3P
     913092-35-0P
                    913092-36-1P
                                   913092-37-2P
                                                                  913092-39-4P
                    913092-41-8P
                                   913092-42-9P
                                                  913092-43-0P
                                                                  913092-44-1P
     913092-40-7P
                    913092-46-3P
                                   913092-47-4P
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     913092-45-2P
                                   913092-76-9P
                                                  913092-77-0P
                                                                  913092-78-1P
     913092-50-9P
                    913092-75-8P
     913976-51-9P, 4-Amino-N-ethyl-3-phenylpiperidine-1-carboxamide
                    913976-59-7P
                                   913976-62-2P
                                                  913976-63-3P
     913976-58-6P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (preparation of piperidine derivs. as tachykinin receptor antagonists)
              THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
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L16 ANSWER 4 OF 9 CAPLUS: COPYRIGHT 2007 ACS on STN
    2005:696924 CAPLUS
ΑN
    143:194181
DN
ED
    Entered STN: 05 Aug 2005
ΤI
    Preparation of novel aminoglycoside antibiotics effective against
    methicillin resistant Staphylococcus aureus (MRSA)
IN
    Minowa, Nobuto; Usui, Takayuki; Akiyama, Yoshihisa; Hiraiwa, Yukiko;
    Yoneda, Toshio; Hasegawa, Toshifumi; Maebashi, Kazunori; Ida, Takashi;
    Katsumata, Kazuko; Otsuka, Keiko; Ikeda, Daishiro
PA
    Meiji Seika Kaisha, Ltd., Japan; Microbial Chemistry Research Foundation
SO
    PCT Int. Appl., 198 pp.
    CODEN: PIXXD2
DT
     Patent
LA
     Japanese
     ICM C07H015-234
IC
     ICS A61K031-7036; A61D031-04
CC
     33-7 (Carbohydrates)
     Section cross-reference(s): 1
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    WO 2005070945 A1
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CLASS
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                        C07H015-234
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                 IPCI
                        CO7HOO15-234 [ICM, 7]; CO7HOO15-00 [ICM, 7, C*];
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A61K0031-7036 [ICS,7]; A61K0031-7028 [ICS,7,C*]; A61D0031-04 [ICS,7]; A61P0031-04 [ICS,7]; A61P0031-00 [ICS, 7, C*] IPCR A61K0031-7028 [I,C*]; A61K0031-7036 [I,A]; C07H0015-00 [I,C*]; C07H0015-234 [I,A] ECLA A61K031/7036; C07H015/234 EP 1710248 IPCI C07H0015-00 [ICM,7] IPCR C07H0015-00 [I,C]; C07H0015-234 [I,A]; A61K0031-7028 [I,C*]; A61K0031-7036 [I,A] ECLA A61K031/7036; C07H015/234

OS MARPAT 143:194181

GΙ

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Arbekacin, 3',4'-dehydroarbekacin, dibekacin, 3',4'-dehydrodibekacin, epiarbekacin, and epidibekacin compds. represented by, e.g., the general formula (I) or pharmacol. acceptable salts or solvates thereof [R4''a, R4''b = H, OH; R5a = halo, HO, NH2, N3, C1-6 alkanoyloxy, C1-6 alkylsulfonyloxy, C1-6 alkanoylamino, arylcarbonylamino, di(C1-6 alkyl)amino, (un)substituted C1-6 alkylamino; R6''a = C1-6 alkyl with ≥1 H atoms optionally being substituted with HO, halo, and NH2; R3''a, R6'a, R6'b = H, C1-6 alkyl; the dotted line is a double or a single bond; m = 0-2; X = H, OH; n = 1-3; * represents R or S configuration; provided that when R4''a = H, R4''b = OH, and the dotted line is a single bond, then R5a = a group described above other than F; or when R3''a = R4''a = H, R6a = hydroxymethyl, R6'a = R6'b = H, and the dotted line = a single bond, then R5a = a described above other than HO, NH2, and N3] are prepared These compds. have excellent antibacterial activity against bacteria causing severe infectious diseases such as pneumonia and sepsis, in particular against MRSA. Also provided are: an antibacterial containing any of these compds.; and a medicinal composition containing any of these compds.

as an active ingredient. Thus, N-protection of arbekacin by reaction with di-tert-Bu dicarbonate in DMF followed by 4'',6''-O-protection by ketalization with 1,1-dimethoxycyclohexane in the presence of p-MeC6H4SO3H.H2O in DMF at 50° and 46-48 mm bar, and O-benzoylation by benzoyl chloride in pyridine at 4-6° for 2 h gave arbekacin derivative (II) which underwent 5-0-mesylation by mesyl chloride in the presence of 4-dimethylaminopyridine in CH2Cl2 at room temperature for 4 h and 5-acetoxylation with cesium acetate in DMF at 100° for 2 h to give 5-epiarbekacin derivative (III). Deketalization of III with 90% aqueous CF3CO2H at room temperature for 4 h, 6''-O-tritylation with trityl chloride in pyridine at 60° for 1 h, 4''-O-triflation with triflic anhydride in the presence of pyridine in CH2Cl2 at -5° for 2 h, and 4''-O-acetoxylation with cesium acetate in DMF at room temperature for 2 h followed by O-deacylation with NaOMe in methanol and then treatment with 90% aqueous CF3CO2H gave 5,4''-diepiarbekacin. The compds. I showed min. inhibitory concentration (MIC) of ≤ 8 , ≤ 4 , and ≤ 4 $\mu g/mL$ against MRSA-HR, MRSA-LR, and Pseudomonas aeruginosa PAO1, resp., while arbekacin showed MIC of 64 and 16 $\mu g/mL$ against MRSA-HR and MRSA-Laser Radiation, resp.

ST aminoglycoside antibiotic prepn antibacterial methicillin resistant Staphylococcus aureus MRSA; arbekacin dehydroarbekacin prepn antibacterial methicillin resistant Staphylococcus aureus; dibekacin dehydrodibekacin prepn antibacterial methicillin resistant Staphylococcus aureus; epiarbekacin epidibekacin prepn antibacterial methicillin resistant

```
Staphylococcus aureus
 ΙT
      Antibiotics
         (aminoglycoside; preparation of aminoglycoside antibiotics effectives as
         antibacterial agents against methicillin resistant Staphylococcus
         aureus (MRSA))
 ΙT
      Infection
         (bacterial; preparation of aminoglycoside antibiotics effectives as
         antibacterial agents against methicillin resistant Staphylococcus
         aureus (MRSA))
 ΙT
      Staphylococcus aureus
         (methicillin-resistant; preparation of aminoglycoside antibiotics effectives
         as antibacterial agents against methicillin resistant Staphylococcus
         aureus (MRSA))
 IT
      Antibacterial agents
         (preparation of aminoglycoside antibiotics effectives as antibacterial
         agents against methicillin resistant Staphylococcus aureus (MRSA))
 ΙT
      Aminoglycosides
      RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
      (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
      (Uses)
         (preparation of aminoglycoside antibiotics effectives as antibacterial
         agents against methicillin resistant Staphylococcus aureus (MRSA))
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      861404-85-5P
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      preparation); THU (Therapeutic use); BIOL (Biological study); PREP
      (Preparation); RACT (Reactant or reagent); USES (Uses)
         (preparation of aminoglycoside antibiotics effectives as antibacterial
         agents against methicillin resistant Staphylococcus aureus (MRSA))
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      (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
      (Uses)
         (preparation of aminoglycoside antibiotics effectives as antibacterial
         agents against methicillin resistant Staphylococcus aureus (MRSA))
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      50-00-0, Formaldehyde, reactions
                                          74-89-5, Methylamine, reactions
      75-07-0, Acetaldehyde, reactions
                                          75-52-5, Nitromethane, reactions
      76-83-5, Triphenylmethyl chloride
                                         77-76-9, 2,2-Dimethoxypropane
      90-02-8, 2-Hydroxybenzaldehyde, reactions
                                                   98-88-4, Benzoyl chloride
      100-39-0, Benzyl bromide
                                 100-52-7, Benzaldehyde, reactions
                                                                      100-83-4,
      m-Hydroxybenzaldehyde
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                                                                  107-11-9,
      Allylamine
                   107-15-3, Ethylenediamine, reactions
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      Acetic anhydride
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                  123-08-0, p-Hydroxybenzaldehyde
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                                           156-87-6, 3-Amino-1-propanol
      141-43-5, 2-Aminoethanol, reactions
                              534-03-2, 2-Amino-1,3-propanediol
      334-88-3, Diazomethane
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                                    925-90-6, Ethylmagnesium bromide
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     26628-22-8, Sodium azide
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     41840-29-3 42491-79-2
                               51025-85-5, Arbekacin
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     N-(tert-Butoxycarbonyl)ethylenediamine
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     N, N'-Bis(tert-butoxycarbonyl)-S-methylisothiourea
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     (Reactant or reagent)
        (preparation of aminoglycoside antibiotics effectives as antibacterial,
        agents against methicillin resistant Staphylococcus aureus (MRSA))
RE.CNT
              THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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     ANSWER 5 OF 9 CAPLUS COPYRIGHT 2007 ACS on STN
L16
     2003:243023 CAPLUS
AN
DN
     139:96917
ED
     Entered STN:
                  30 Mar 2003
     Structural regulation of a peptide-conjugated graft copolymer: A simple
TΙ
     model for amyloid formation
ΑU
     Koga, Tomoyuki; Taguchi, Kazuhiro; Kobuke, Yoshiaki; Kinoshita, Takatoshi;
     Higuchi, Masahiro
     Nanoarchitectonics Research Center, National Institute of Advanced
CS
     Industrial Science and Technology and CREST (Japan Science and
     Technology), Ibaraki, 305-8565, Japan
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SO
     Chemistry--A European Journal (2003), 9(5), 1146-1156
     CODEN: CEUJED; ISSN: 0947-6539
PB
     Wiley-VCH Verlag GmbH & Co. KGaA
DΤ
     Journal
LA
     English
CC
     6-3 (General Biochemistry)
     CASREACT 139:96917
OS
AB
     The self-assembly of peptides and proteins into \beta-sheet-rich
     high-order structures has attracted much attention as a result of the
     characteristic nanostructure of these assemblies and because of their
     association with neurodegenerative diseases. Here we report the structural
     and conformational properties of a peptide-conjugated graft copolymer,
     poly(\gamma-methyl-L-glutamate) grafted polyallylamine (1) in a
     water-2,2,2-trifluoroethanol solution as a simple model for amyloid
     formation. Atomic force microscopy revealed that the globular peptide 1
     self-assembles into nonbranching fibrils that are about 4 nm in height
     under certain conditions. These fibrils are rich in \beta-sheets and,
     similar to authentic amyloid fibrils, bind the amyloidophilic dye Congo
     red. The secondary and quaternary structures of the peptide 1 can be
     controlled by manipulating the pH, solution composition, and salt
concentration; this
     indicates that the three-dimensional packing arrangement of peptide chains
     is the key factor for such fibril formation. Furthermore, the addition of
     carboxylic acid-terminated poly(ethylene glycol), which interacts with
     both of amino groups of 1 and hydrophobic PMLG chains, was found to
     obviously inhibit the \alpha-to-\beta structural transition for
     non-assembled peptide 1 and to partially cause a \beta-to-\alpha
     structural transition against the 1-assembly in the \beta-sheet form.
     These findings demonstrate that the amyloid fibril formation is not
     restricted to specific protein sequences but rather is a generic property
     of peptides. The ability to control the assembled structure of the
     peptide should provide useful information not only for understanding the
     amyloid fibril formation, but also for developing novel peptide-based
     material with well-defined nanostructures.
ST
     peptide conjugated graft copolymer amyloid fibril model
ΙT
     Fibril
        (amyloid-like; peptide-conjugated graft copolymer as a simple model for
        amyloid fibril formation)
ΙT
        (fibril, amyloid-like; peptide-conjugated graft copolymer as a simple
        model for amyloid fibril formation)
IΤ
     Polymers, biological studies
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (graft; peptide-conjugated graft copolymer as a simple model for
        amyloid fibril formation)
ΙT
     Self-assembly
        (into amyloid-like fibrils; peptide-conjugated graft copolymer as a
        simple model for amyloid fibril formation)
     Conformation
     β-Sheet
        (peptide-conjugated graft copolymer as a simple model for amyloid
        fibril formation)
ΙT
     Amyloid
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (peptide-conjugated graft copolymer as a simple model for amyloid
        fibril formation)
     36877-69-7DP, reaction products with allylamine-Me glutamate
ΙT
     graft copolymer/nitrobenzofurazan derivs.
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP
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(Preparation); USES (Uses)
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        peptide-conjugated graft copolymer as a simple model for amyloid fibril
        formation)
ΙT
     29270-56-2DP, reaction products with allylamine-Me glutamate
     graft copolymer
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
         (nitrobenzofurazan-containing peptide-conjugated graft copolymer;
        peptide-conjugated graft copolymer as a simple model for amyloid fibril
        formation)
IT
     1663-47-4, \gamma-Methyl-L-glutamate-N-carboxy anhydride
     24424-99-5, Di-tert-butyl dicarbonate
                                              29270-56-2,
     4-Fluoro-7-nitrobenzofurazan
                                     30551-89-4, Polyallylamine
                                                                  36877-69-7,
     Rhodamine B isothiocyanate
     RL: RCT (Reactant); RACT (Reactant or reagent)
         (peptide-conjugated graft copolymer as a simple model for amyloid
        fibril formation)
ΙT
     431047-85-7DP, BOC-protected, benzofurazan derivs., reaction products with
     Rhodamine B isothiocyanate
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
         (peptide-conjugated graft copolymer; peptide-conjugated graft copolymer
        as a simple model for amyloid fibril formation)
RE.CNT
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     Synthesis of 1,3-diamino-4-(aminomethyl)-benzene derivates and their use
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IN
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     Wella AG, Germany
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     Ger. Offen., 16 pp.
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          C07C215-08; C07C217-00; C07C211-52; C07C211-53; D06P001-32;
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ECLA
                       C07C211/51; C07C215/14; C07C217/08; C07C233/36;
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os
    MARPAT 137:252674
AΒ
    The invention concerns the synthesis of 1,3-diamino-4-(aminomethyl)-
    benzene derivates and their use as coupling agents in oxidative hair dyes.
    The hair prepns. further contain developers, other coupling agents and
    direct dyes. Thus 1,3-diamino-4-(methylaminomethyl)-benzene hydrochloride
    was synthesized and used as a 1.25 mmol coupler ingredient in a hair dye
    that contained 1.25 mmol 1,4-diamino benzene as developer. Further
    ingredients were (g); potassium oleate (8% aqueous solution) 1.0;
    ammonia (22% aqueous solution) 1.0; ethanol 1.0; ascorbic acid 0.3;
    water to 100.
ST
    diamino aminomethyl benzene derivate oxidative hair dye
ΙT
    Dyes
        (direct; synthesis of 1,3-diamino-4-(aminomethyl)-benzene derivates and
       use in oxidative hair dyes)
TT
    Hair preparations
        (dyes, oxidative; synthesis of 1,3-diamino-4-(aminomethyl)-benzene
        derivates and use in oxidative hair dyes)
ΙT
                   460990-07-2P
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        (coupling agent; synthesis of 1,3-diamino-4-(aminomethyl)-benzene
        derivates and use in oxidative hair dyes)
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     1,7-Dihydroxynaphthalene
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     110-91-8, Morpholine, reactions
                                     123-75-1, Pyrrolidine, reactions
     141-43-5, Ethanolamine, reactions
                                        498-63-5, Prolinol
                                                              1001-53-2,
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                                 4795-29-3, Tetrahydrofurfuryl amine
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                                      6168-72-5, 2-Aminopropanol
                                                                   6859-99-0,
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     40499-83-0, 3-Pyrrolidinol
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IT
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L16
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ED
TΙ
     Towards self-assembled electro- and photo-active organic nanotubes
ΑU
     Fenniri, Hicham; Packiarajan, Mathivanan; Ribbe, Alexander E.; Vidale,
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CS
     1393 H. C. Brown Laboratory of Chemistry, Purdue University, West
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     Polymer Preprints (American Chemical Society, Division of Polymer
SO
     Chemistry) (2001), 42(2), 569-670
     CODEN: ACPPAY; ISSN: 0032-3934
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DT
     Journal; (computer optical disk)
LA
     English
     73-11 (Optical, Electron, and Mass Spectroscopy and Other Related
CC
     Properties)
     Section cross-reference(s): 22, 76
AΒ
     The synthesis, self-assembly, and characterization of the organic nanotubular
     scaffolds from self-assembled supermacrocycles of low-mol. weight synthetic
     modules, and its scope in the design of 1 dimensional, reversible,
     polymeric materials with photonic and electronic properties are discussed.
     A hybrid self-complementary base (M1) composed of the Watson-Crick
     donor-donor-acceptor hydrogen bonding array of guanine and the
     acceptor-acceptor-donor array of cytosine was synthesized. Electrospray
     ionization mass spectrometry of dilute aqueous solns. of M1 showed all
     the peaks corresponding to the noncovalent intermediate species of the
     parent rosette. M1 underwent a cooperative, hierarchical self-assembly
     process through H-bonding, stacking interactions, and hydrophobic effects.
     The system established that electrostatic, stacking, and hydrophobic
     interactions could be effectively orchestrated by hydrogen bonds to direct
     the hierarchical assembly and organization of helical nanotubular
     architectures in an aqueous milieu. The amenability of the
     synthetic scheme to oligomerization and incorporation of alternative
     components and bases will provide a rapid access to tubular assemblies
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with predefined dimensions, photonic, electronic, and transport

(of self-assembled electro- and photo-active organic nanotubes)

nanotube self assembly supermacrocycle Watson Crick bonding

ST IT properties.

Circular dichroism

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·IT
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RE.CNT
               THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
(1) Chopra, N; Science 1995, V269, P966 CAPLUS
(2) Fenniri, H; J Am Chem Soc 2001, V123, P3854 CAPLUS
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     Hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivatives
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     Chassot, Laurent; Baun, Hans-Jurgen
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     Wella Aktiengesellschaft, Germany
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     Eur. Pat. Appl., 31 pp.
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                       C07C233/36; C07C237/06; C07C239/20; C07C255/03;
                       C07C311/37; C07C323/58; C07D295/12A2; C07D295/12A3;
                       C07D295/12B1B1
JP 2001199941
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                       C07C0211-50 [ICS,7]; C07C0211-00 [ICS,7,C*];
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OS MARPAT 135:111699

The invention concerns oxidative hair dyes that contain as developers AB 2-aminoalkyl-1,4-diaminobenzene derivs. or their physiol. compatible water soluble salts of the formula (I), where R1-R7 are defined. The hair dye compns. further contain another developer, e.g. 1,4-diamiaminobenzene, 2,5-diaminotoluene; coupling agents, e.g. 2,6-diaminopyridine; and at least one direct dye. Thus, bromo-p-phenylenediamine-HCl was converted with di-tert-Bu dicarbonate to 2,5-bis(tert-butoxycarbonylamino)bromobenzene, and then with DMF in the presence of methyllithium and butyllithium to (2-formyl-1,4phenylene)biscarbamic acid di(tert-butyl)ester. This compound was reacted with ethylamine and the hydrochloride of the formed substance was prepared The obtained 2-ethylaminomethy-1,4-diamino benzene hydrochloride was used (0.0125 mmol) in a hair dye, that further contained: 1,3-dihydroxybenzene (coupling agent) 0.0125 mmol; potassium oleate (8% aqueous solution) 0.01 g; ammonia (22% aqueous solution) 0.01 g; ethanol 0.01 g; ascorbic acid 0.003 g; water to 1 g. The dye resulted a light blond color.

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ST
     aminoalkyl diaminobenzene deriv oxidative hair dye
ΙT
     Dyes
        (direct; hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivs.)
ΙT
     Hair preparations
        (dyes, oxidative; hair dyes containing 2-aminoalkyl-1,4-diaminobenzene
        derivs.)
IT
     Coupling agents
        (hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivs.)
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (2,5-Diaminotoluene sulfate; hair dyes containing 2-aminoalky1-1,4-
        diaminobenzene derivs.)
IT
     6358-09-4, 2-Amino-6-chloro-4-nitrophenol
                                                  28365-08-4
                                                               53347~10-7
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (direct dye; hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivs.)
ΙT
     83-56-7, 1,5-Dihydroxynaphthalene
                                        89-25-8, 3-Methyl-1-phenyl-5-
                  89-83-8, 5-Methyl-2-(1-methylethyl)phenol 90-15-3, 91-56-5, 2,3-Indolinedione 91-68-9, 3-Diethylaminophenol
     pyrazolone
     1-Naphthol
     92-44-4, 2,3-Dihydroxynaphthalene
                                         95-70-5, 2,5-Diaminotoluene
                                                                        95-88-5,
     1,3-Benzenediol, 4-chloro- 99-07-0, 3-Dimethylaminophenol
                                                                    106-50-3,
     1,4-Diaminobenzene, biological studies
                                               106-50-3D, 1,4-Benzenediamine,
     2-aminoalkyl derivs., biological studies
                                                108-45-2, 1,3-Diaminobenzene,
     biological studies 108-46-3, 1,3-Dihydroxybenzene, biological studies
                141-86-6, 2,6-Diaminopyridine
     137-19-9
                                                533-31-3, 3,4-
                           575-38-2, 1,7-Dihydroxynaphthalene
     Methylenedioxyphenol
                                                                  582-17-2,
     2,7-Dihydroxynaphthalene
                                608-25-3, 1,3-Dihydroxy-2-methylbenzene
     619-05-6, 3,4-Diaminobenzoic acid
                                         770-25-2
                                                    1953-54-4, 5-Hydroxyindole
     2380-84-9, 7-Hydroxyindole 2380-86-1, 6-Hydroxyindole 4-Hydroxyindole 2835-99-6, 3-Methyl-4-aminophenol 31
                                                                2380-94-1,
                                                           3131-52-0,
     5,6-Dihydroxyindole 5349-76-8, 2,4-Diamino-1-methoxy-5-methylbenzene
     5697-02-9, 2-Methyl-1-naphthyl-acetate
                                              6201-65-6, 2-Chloro-1,3-
     dihydroxybenzene 6265-21-0, 3-[(2-Hydroxyethyl)amino]aniline
     6941-70-4, 6-Bromo-1-hydroxy-3,4-methylenedioxybenzene
                                                               7228-00-4
     16867-03-1, 2-Amino-3-hydroxypyridine 26011-57-4, 6-Amino-3,4-dihydro-
     1,4(2H)benzoxazine 26021-57-8, 3,4-Dihydro-6-hydroxy-1,4(2H)benzoxazine
     26455-21-0, N-(3-Dimethylamino)phenylurea
                                                  28020-38-4,
     2,3-Diamino-6-methoxypyridine 29539-03-5, 5,6-Dihydroxyindoline
     39489-79-7, 5-Amino-2, 4-dichloro-phenol 53222-92-7, 3-Amino-2-
     methylphenol
                    54381-16-7
                                 55302-96-0, 5-[(2-Hydroxyethyl)amino]-2-
     methylphenol
                    56216-28-5, 3,5-Diamino-2,6-dimethoxypyridine-
                       61693-42-3, 3-Amino-2,4-dichloro-phenol
                                                                  70643-19-5,
     dihydrochloride
     2,4-Diamino-1-(2-hydroxyethoxy)benzene 70643-20-8
                                                           71500-41-9
     71500-42-0
                  74918-21-1, 1,3-Bis(2,4-Diaminophenoxy)propane-
     tetrahydrochloride
                          76045-64-2
                                        78661-33-3
                                                     80592-80-9 80592-81-0
                              83763-47-7, 2-Amino-4-[(2-
     81329-90-0
                  81892-72-0
                                  83763-48-8
     hydroxyethyl)amino]anisole
                                                84540-48-7
                                                             84540-50-1,
     3-Amino-2-chloro-6-methylphenol
                                        86817-42-7, 2-(4-Amino-2-
     hydroxyphenoxy) ethanol
                              90817-34-8, 3-Amino-6-methoxy-2-
     (methylamino)pyridine
                             93841-24-8, 2-(2,5-Diaminophenyl)ethanol
                  94082-77-6 104752-50-3
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     93841-25-9
                                                            110102-86-8,
     5-Amino-4-chloro-2-methylphenol
                                       111451-24-2, 2,6-Diamino-3,5-
     dimethoxypyridine
                         115423-86-4, 1,3-Diamino-2,4-dimethoxybenzene
     122455-85-0
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                       137290-78-9, 5-Amino-4-methoxy-2-methylphenol
     dihydrochloride
     139443-57-5, 5-Amino-4-ethoxy-2-methylphenol
                                                    141614-04-2,
     1,3-Benzenediamine, 4-ethoxy-6-methyl- 141614-05-3, 2,4-Diamino-1-(2-
     hydroxyethoxy) -5-methylbenzene
                                       141922-20-5, 2,4-Diamino-1-fluoro-5-
                     142082-56-2
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     methylbenzene
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                  207923-07-7
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     168092-23-7
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
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```
(hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivs.)
ΙT
     123-30-8, 4-Aminophenol
                             591-27-5, 3-Aminophenol
     RL: BSU (Biological study, unclassified); RCT (Reactant); BIOL (Biological
     study); RACT (Reactant or reagent)
        (hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivs.)
IT
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     RL: BSU (Biological study, unclassified); SPN (Synthetic preparation);
     BIOL (Biological study); PREP (Preparation)
        (hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivs.)
IT
     59-51-8, Methionine
                          62-53-3, Aniline, reactions 75-04-7, Ethylamine,
                 75-31-0, Isopropylamine, reactions
                                                      95-85-2,
     reactions
                              97-51-8, 2-Hydroxy-5-nitrobenzaldehyde
     4-Chloro-2-aminophenol
                                                                        98-03-3,
                                99-57-0, 2-Amino-4-nitrophenol
                                                                  99-98-9,
     Thiophene-2-carbaldehyde
     4-Amino-N, N-dimethylaniline 100-52-7, Benzaldehyde, reactions
                                    106-47-8, 4-Chloroaniline, reactions
     104-86-9, 4-Chlorobenzylamine
     106-49-0, 4-Methylaniline, reactions
                                           107-10-8, Propylamine, reactions
                            108-00-9, 2-Dimethylamino ethylamine
     107-11-9, Allylamine
     109-01-3
               109-55-7, 3-Dimethylamino propylamine
                                                        109-83-1, 2-Methylamino
               109-85-3, 2-Methoxy ethylamine 110-58-7, Pentylamine
     110-73-6, 2-Ethylamino-ethanol 110-91-8, Morpholine, reactions
                                          120-57-0, 3,4-
     111-42-2, Diethanolamine, reactions
     Methylenedioxybenzaldehyde
                                  123-08-0, 4-Hydroxybenzaldehyde
                                                                     123-72-8,
     Butyraldehyde
                    123-75-1, Pyrrolidine, reactions
                                                        364-73-8, Benzene,
     4-bromo-1-fluoro-2-nitro-
                                364-74-9, 1,4-Difluoro-2-nitrobenzene
               437-83-2, 3-Fluoro-2-methoxy aniline 446-35-5,
     1,3-Difluoro-4-nitrobenzene 455-14-1, 4-Trifluoromethyl aniline
     498-63-5, Prolinol
                          500-22-1, Pyridine-3-carbaldehyde 536-21-0,
     1-(3-Hydroxyphenyl)-2-aminoethanol
                                          536-90-3, 3-Methoxyaniline
     555-16-8, 4-Nitrobenzaldehyde, reactions
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     Dimethylaminobenzaldehyde 587-04-2, 3-Chlorobenzaldehyde
                                                                   590-86-3,
     3-Methylbutyraldehyde 609-36-9, Proline 616-30-8, 3-Amino-1,2-propane
            617-45-8, Aspartic acid 617-89-0, Furfurylamine
                        872-85-5, Pyridine-4-carbaldehyde
                                                             1117-97-1,
     Cyclopropylamine
     O, N-Dimethyl-hydroxylamine 1121-60-4, 2-Pyridinecarboxaldehyde
     1493-27-2, 1-Fluoro-2-nitrobenzene
                                          2038-03-1, 4-Morpholineethanamine
     2043-61-0, Cyclohexane carbaldehyde 2454-37-7, 3-(1-Hydroxyethyl)-aniline 2516-47-4, Aminomethyl cyclopropane 2812-47-7, Prolinamide
     2835-95-2, 3-Amino-6-methylphenol 3731-51-9, 2-Picolylamine
                     4214-76-0, 2-Amino-5-nitropyridine
                                                            4795-29-3,
     4-Picolylamine
     Tetrahydrofurfurylamine 5036-48-6, 1-(3-Aminopropyl)imidazole
     5382-16-1, 4-Hydroxypiperidine 5616-32-0, Methylaminoacetonitrile
     6168-72-5, 2-Aminopropanol
                                  6291-85-6, 3-Ethoxypropylamine
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3,4-Dimethoxy aniline 6859-99-0, 3-Hydroxypiperidine 6921-22-8
    7304-32-7, 2-Fluoro-5-nitro benzoic acid 7663-77-6, 1-(3-Aminopropyl)-2-
                                              14268-66-7, 3,4-Methylene
    pyrrolidone
                 13325-10-5, 4-Aminobutanol
    dioxyaniline 24424-99-5, Di-tert-butyl dicarbonate 25739-59-7
     35303-76-5, 4-(2-Aminoethyl)-benzenesulfonamide 40499-83-0,
     3-Hydroxypyrrolidine 51980-54-2, 4-Pyrrolidino benzaldehyde 68621-88-5
     71026-66-9 244104-66-3
                               325953-40-0
                                           325953-41-1 325953-45-5
                 325953-48-8
     325953-46-6
    RL: RCT (Reactant); RACT (Reactant or reagent)
        (hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivs.)
                   325953-36-4P
                                 350481-97-9P
IΤ
    244104-65-2P
                                                350481-98-0P
    RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (hair dyes containing 2-aminoalkyl-1,4-diaminobenzene derivs.)
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AN
    1996:353642 CAPLUS
    125:34158
DN
ΕD
    Entered STN: 19 Jun 1996
ŤΙ
    Preparation of N-[5-[2-(2-amino-4-oxo-4,6,7,8-tetrahydro-3H-pyrimido[5,4-
    b][1,4]thiazin-6-y1)ethy1]-2-thienoy1]-L-glutamic acid derivative and
     analog as antiproliferative agents and glycinamide ribonucleotide formyl
     transferase (GARFT) inhibitors
    Varney, Michael D.; Romines, William H.; Palmer, Cynthia L.
IN
PA
    Agouron Pharmaceuticals, Inc., USA
SO
     PCT Int. Appl., 58 pp.
    CODEN: PIXXD2
DT
     Patent
LA
     English
     ICM C07D513-04
IC
     ICS C07D498-04; C07D517-04; C07D333-38; C07D417-06; A61K031-54
     C07D513-04, C07D279-00, C07D239-00; C07D498-04, C07D265-00, C07D239-00;
     C07D517-04, C07D293-00, C07D239-00
CC
     34-2 (Amino Acids, Peptides, and Proteins)
     Section cross-reference(s): 1
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                              DATE
                                          APPLICATION NO.
     PATENT NO.
                        KIND
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                        A1 19960208 WO 1994-US8522
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            NL, NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TJ, TT, UA, UZ, VN
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19940728 WO .1994-US8522 CLASS PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES WO 9603407 ICM C07D513-04 ICS C07D498-04; C07D517-04; C07D333-38; C07D417-06; A61K031-54 C07D513-04, C07D279-00, C07D239-00; C07D498-04, C07D265-00, C07D239-00; C07D517-04, C07D293-00, ICI C07D239-00 IPCI C07D0513-04 [ICM, 6]; C07D0498-04 [ICS, 6]; C07D0517-04 [ICS, 6]; C07D0333-38 [ICS, 6]; C07D0333-00 [ICS, 6, C*]; C07D0417-06 [ICS, 6]; C07D0417-00 [ICS, 6, C*]; A61K0031-54 [ICS, 6]; C07D0513-04 [ICI, 6]; C07D0513-00 [ICI, 6, C*]; C07D0279-00 [ICI, 6]; C07D0239-00 [ICI, 6]; C07D0498-04 [ICI,6]; C07D0498-00 [ICI,6,C*]; C07D0265-00 [ICI,6]; C07D0239-00 [ICI,6]; C07D0517-04 [ICI, 6]; C07D0517-00 [ICI, 6, C*]; C07D0293-00 [ICI, 6]; C07D0239-00 [ICI,6] IPCR C07D0333-00 [I,C*]; C07D0333-38 [I,A]; C07D0417-00 [I,C*]; C07D0417-06 [I,A]; C07D0498-00 [I,C*]; C07D0498-04 [I,A]; C07D0513-00 [I,C*]; C07D0513-04 [I,A]; C07D0517-00 [I,C*]; C07D0517-04 [I,A] ECLA C07D333/38; C07D417/06+333B+279; C07D498/04+265B+239B; C07D513/04+279B+239B; C07D517/04+293B+239B AU 9475520 IPCI C07D0513-04 [ICM, 6]; C07D0513-00 [ICM, 6, C*]; C07D0498-04 [ICS,6]; C07D0498-00 [ICS,6,C*]; C07D0517-04 [ICS,6]; C07D0517-00 [ICS,6,C*]; C07D0333-38 [ICS, 6]; C07D0333-00 [ICS, 6, C*]; C07D0417-06 [ICS, 6]; C07D0417-00 [ICS, 6, C*]; A61K0031-54 [ICS, 6] IPCR A61K [I,S]; A61K0031-54 [I,C*]; A61K0031-54 [I,A]; C07D [I,S]; C07D0333-00 [I,C*]; C07D0333-38 [I,A]; C07D0417-00 [I,C*]; C07D0417-06 [I,A]; C07D0498-00 [I,C*]; C07D0498-04 [I,A]; C07D0513-00 [I,C*]; C07D0513-04 [I,A]; C07D0517-00 [I,C*]; C07D0517-04 EP 783507 IPCI C07D0513-04 [ICM, 6]; C07D0498-04 [ICS, 6]; C07D0517-04 [ICS, 6]; C07D0333-38 [ICS, 6]; C07D0333-00 [ICS, 6, C*]; C07D0417-06 [ICS, 6]; C07D0417-00 [ICS, 6, C*]; A61K0031-54 [ICS,6]; C07D0513-04 [ICI,6]; C07D0513-00 [ICI, 6, C*]; C07D0279-00 [ICI, 6]; C07D0239-00 [ICI, 6]; C07D0498-04 [ICI,6]; C07D0498-00 [ICI,6,C*]; C07D0265-00 [ICI,6]; C07D0239-00 [ICI,6]; C07D0517-04 [ICI, 6]; C07D0517-00 [ICI, 6, C*]; C07D0293-00 [ICI, 6]; . C07D0239-00 [ICI,6] IPCR A61K [I,S]; A61K0031-54 [I,C*]; A61K0031-54 [I,A]; C07D [I,S]; C07D0333-00 [I,C*]; C07D0333-38 [I,A]; C07D0417-00 [I,C*]; C07D0417-06 [I,A]; C07D0498-00 [I,C*]; C07D0498-04 [I,A]; C07D0513-00 [I,C*]; C07D0513-04 [I,A]; C07D0517-00 [I,C*]; C07D0517-04 [I,A]JP 10504541 IPCI C07D0498-04; C07D0498-00 [C*]; A61K0031-505; C07D0513-04; C07D0513-00 [C*]; C07D0517-04; C07D0517-00 [C*] RU 2136686 IPCI C07D0513-04 [ICM, 6]; C07D0417-06 [ICS, 6]; C07D0333-38 [ICS, 6]; C07D0333-40 [ICS, 6]; A61K0031-54 [ICS, 6]; C07D0513-04 [ICI,6]; C07D0513-00 [ICI,6,C*]; C07D0239-00 [ICI,6]; C07D0279-14 [ICI,6]; C07D0417-06 [ICI,6]; C07D0417-00 [ICI,6,C*]; C07D0279-12 [ICI,6];

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OS
     MARPAT 125:34158
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OS MARPAT 125:34158 GI

The title compds. [I; A = O, S, Se; X = each (un) substituted C1-3 alkyl, AΒ C2-3 alkenyl, C2-3 alkynyl, or amino group, S, O; Y = O, S, NH; B = H, halo; C = H, halo, (un) substituted C1-6 alkyl; R1, R2 = H, a moiety that forms with the attached CO2 a readily hydrolyzable ester group] in equilibrium with its 4-hydroxy tautomer, and its pharmaceutically acceptable salts are prepared Thus, 5-[2-(2-amino-4-oxo-4,6,7,8-tetrahydro-3H-pyrimidino[5,4b][1,4]thiazin-6-yl)ethyl]-2-thienoic acid (preparation given) was condensed with H-Glu(OEt)-OEt.HCl using 1-hydroxybenzotriazole hydrate, (Me2CH)2NEt, and 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride in DMF at 25° for 20 h to give the title compound di-Et ester I (A = Y = S, B = C = H, X = CH2, R1 = R2 = Me) (71%), which was stirred at 25° for 14 h, cooled to 0°, and adjusted to pH 3.5 with aqueous HCl to give 80% the title compound I (A = Y = S, B = C = R1 = R2 = H, X = CH2). The latter compound in vitro showed competitive inhibition of human GARFT with Ki value of 4.5 nM and IC50 of 16 and 4.3 nM for inhibiting the proliferation of L1210 murine leukemia and CCRF-CEM leukemia cells, resp. It at 12.5 mg/kg s.c. in vivo inhibited 100% the growth of lymphosarcoma 6C3HED tumor implanted in C3H/He female mice. ST aminooxotetrahydropyrimidothiazinylethylthienoylglutamic acid prepn antitumor; pyrimidothiazinylethylthienoylglutamic acid prepn antitumor; thienoylglutamic acid pyrimidothiazinylethyl prepn antitumor; glutamic acid pyrimidothiazinylethyl thienoyl prepn antitumor; glycinamide ribonucleotide formyl transferase GARFT inhibitor IT Neoplasm inhibitors

Ι

(preparation of N-[[(aminooxotetrahydropyrimidothiazinyl)ethyl]thienoyl]glut amic acid derivative and analog as antitumor agents and glycinamide

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ribonucleotide formyl transferase inhibitors)
ΙT
     177575-16-5P
                    177575-17-6P
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                                                  177575-19-8P
                                                                 177575-20-1P
     177575-21-2P
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     BIOL (Biological study); PREP (Preparation); USES (Uses)
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        ribonucleotide formyl transferase inhibitors)
ΙT
     9032-02-4
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     (Miscellaneous); BIOL (Biological study); PROC (Process)
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        ribonucleotide formyl transferase inhibitors)
IT
     50-01-1, Guanidine hydrochloride 107-11-9, Allylamine
     1066-54-2, Trimethylsilylacetylene 1118-89-4, L-Glutamic acid diethyl
     ester hydrochloride
                          10387-40-3, Potassium thioacetate 24424-99-5
                                  26690-80-2
                                               28868-76-0, Dimethyl
     , Di-tert-butyl dicarbonate
     chloromalonate
     RL: RCT (Reactant); RACT (Reactant or reagent)
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        ribonucleotide formyl transferase inhibitors)
TΤ
     62224-19-5P, Methyl 5-bromo-2-thiophenecarboxylate
                                                          89711-08-0P
     115269-99-3P, N, N-Bis(tert-butoxycarbonyl)allylamine
     155089-06-8P, N,N-Bis(tert-butoxycarbonyl)glycinal
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     177575-27-8P
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     177575-32-5P
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     (Reactant or reagent)
        (preparation of N-[[(aminooxotetrahydropyrimidothiazinyl)ethyl]thienoyl]glut
        amic acid derivative and analog as antitumor agents and glycinamide
        ribonucleotide formyl transferase inhibitors)
=> d his
     (FILE 'HOME' ENTERED AT 18:44:27 ON 26 JUN 2007)
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L1
           1332 S ALLYLAMINE
L2
            505 S L1 AND POLY
L3
              0 S METHYLCARBOXY AND L2
     FILE 'CAPLUS' ENTERED AT 18:45:45 ON 26 JUN 2007
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L4
            109 S L4 AND ALLYL
L5
L6
              3 S L5 AND POLY
L7
           1966 S POLYALLYLAMINE
             19 S L7 AND PROTECTED
L8
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                SET NOTICE LOGIN DISPLAY
L10
              0 S POLYALLYLAMINE AND L9
              0 S AMINE AND L9
L11
              1 S L9
L12
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11/245136

CA SUBSCRIBER PRICE

FILE 'CAPLUS' ENTERED AT 18:51:28 ON 26 JUN 2007

L13 5391 S L9
L14 76 S L9 AND ALLYLAMINE
L15 6 S L14 AND WATER
L16 9 S L14 AND (AQ OR WATER)

=> log y
COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION
FULL ESTIMATED COST 37.51 151.33

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

SESSION

-7.02 -24.18

ENTRY

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